IDENTIFICATION OF SORTASE GENE

GOVERNMENT RIGHTS

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BACKGROUND OF THE INVENTION

This invention is directed to an enzyme from Gram-positive bacteria, designated sortase-transamidase, nucleic acid segments encoding the enzyme, and methods of use of the enzyme.

Human infections caused by Gram-positive bacteria present a medical challenge due to the dramatic increase in multiple antibiotic resistance strains in recent years. Gram-positive bacteria that can cause serious or fatal infections in humans include *Staphylococcus*, *Streptococcus*, *Enterococcus*, *Pneumococcus*, *Bacillus*, *Actinomyces*, *Mycobacterium*, and *Listeria*, as well as others. Infections caused by these pathogens are particularly severe and difficult to treat in immunologically compromised patients. These include patients suffering from infection with the Human Immunodeficiency Virus (HIV), the virus that causes AIDS, as well as patients given immune-suppressive agents for treatment of cancer or autoimmune diseases. In particular, infections caused by various *Mycobacterium* species, including *M. tuberculosis*, *M. bovis*, *M. avium*, and *M. intracellulare*, are frequently the cause of disease in patients with AIDS.

Therefore, it is apparent that new target sites for bacterial chemotherapy are needed if such pathogenic organisms are to be controlled.

A unique characteristic of these pathogens and many Gram-positive bacteria is their surface display of proteins anchored to the cell wall. In fact, many of these molecules are known to be involved in essential cellular functions, including pathogenesis in a susceptible host.

Thus, a possible disruption in this anchoring process may prove to be an effective treatment against these disease-causing elements.

The anchoring of surface molecules to the cell wall in Gram-positive bacteria has been demonstrated to involve a conserved pathway, culminating in recognition of a conserved cleavage/anchoring site by some previously uncharacterized cellular machinery. Molecules whose ultimate location is the cell wall must invariably be translocated across the single cellular membrane of these organisms. This is mediated for all cell wall anchored proteins by the well studied secretory pathway, involving cleavage of an amino-terminal signal peptide by a type I signal peptidase. Upon translocation of the molecule out of the cytoplasm, a mechanism must be present that extracellularly recognizes this protein as a substrate for anchoring. This process has been previously shown to involve the carboxyl-terminally located cell wall sorting signal, consisting of a highly conserved motif such as LPXTG (SEQ ID NO:1), in which X can represent any of the twenty naturally occurring L-amino acids, followed by a series of hydrophobic residues and ultimately a sequence of positively-charged residues. Thus, once amino-terminally modified and successfully secreted, a polypeptide with this carboxyl-terminal sequence can present itself as a substrate to be processed by the anchoring machinery. At this time, cleavage of the sorting signal after the threonine residue is coupled with covalent linkage of the remainder of the polypeptide to the free amino group of the pentaglycine crossbridge in the cell wall.

It is this transpeptidation reaction that anchors mature surface proteins to the peptidoglycan layer, from which point the molecules can serve their biological functions. Therefore, there is a need to isolate and purify the enzyme that catalyzes this reaction. There is also a need to identify the gene encoding such an enzyme in order that the enzyme can be produced by genetic engineering techniques. There is also a need to identify compounds that interfere with surface protein anchoring by inhibiting sortase.

Additionally, there is also a need to develop new methods for displaying proteins or peptides on the surfaces of bacteria. For many purposes, it is desirable to display proteins or peptides on the surfaces of bacteria so that the proteins or peptides are accessible to the

surrounding solution, and can, for example, be bound by a ligand that is bound specifically by the protein or peptide. In particular, the display of proteins on the surface of bacteria is desirable for the preparation of vaccines, the linkage of molecules such as antibiotic molecules or diagnostic reagents to cells, for screening reagents such as monoclonal antibodies, and for the selection of cloned proteins by displaying the cloned proteins, then observing their reaction with specific reagents such as antibodies. One way of doing this has been with phage display (G.P. Smith, "Filamentous Fusion Phage: Novel Expression Vectors that Display Cloned Antigens on the Virion Surface," Science 228:1315-1316 (1985)). However, phage display is limited in its practicality, because it requires that the protein being displayed to be inserted into a coat protein of filamentous phage and retain its activity while not distorting the conformation of the coat protein, allowing functional virions to be formed. In general, this technique is therefore limited only to small peptide and proteins.

Therefore, there is a need for a more general method of peptide and protein display.

SUMMARY

The present invention is directed to sortase-transamidase enzymes from Gram-positive bacteria, particularly the products of the surface protein sorting (*srt*A) gene of *Staphylococcus aureus*, and methods for their use, particularly in the areas of drug screening and peptide and protein display.

One aspect of the present invention is a substantially purified sortase-transamidase enzyme from a Gram-positive bacterium, the enzyme catalyzing a reaction that covalently cross-links the carboxyl terminus of a protein having a sorting signal to the peptidoglycan of a Gram-positive bacterium, the sorting signal having a motif of LPX_3X_4G therein, wherein sorting occurs by cleavage between the fourth and fifth residues of the LPX_3X_4G motif. Typically, the Gram-positive bacterium is a species selected from the group consisting of but not limited to *Staphylococcus aureus*, *S. sobrinus*, *Enterococcus faecalis*, *Streptococcus pyogenes*, and *Listeria*

monocytogenes. Preferably, the Gram-positive bacterium is *S. aureus*, and more preferably, the enzyme is the product of the *srt*A gene of *S. aureus*.

Preferably, the enzyme has a molecular weight of about 23,539 daltons and the sorting signal further includes: (2) a substantially hydrophobic domain of at least 31 amino acids carboxyl to the motif; and (3) a charged tail region with at least two positively charged residues carboxyl to the substantially hydrophobic domain, at least one of the two positively charged residues being arginine, the two positively charged residues being located at residues 31-33 from the motif, wherein X_3 is any of the twenty naturally-occurring L-amino acids and X_4 is selected from the group consisting of alanine, serine, and threonine.

Preferably, the enzyme includes an amino acid sequence of: (1) M-K-K-W-T-N-R-L-M-T-I-A-G-V-V-L-I-L-V-A-A-Y-L-F-A-K-P-H-I-D-N-Y-L-H-D-K-D-K-D-E-K-I-E-Q-Y-D-K-N-V-K-E-Q-A-S-K-D-K-K-Q-Q-A-K-P-Q-I-P-K-D-K-S-K-V-A-G-Y-I-E-I-P-D-A-D-I-K-E-P-V-Y-P-G-P-A-T-P-E-Q-L-N-R-G-V-S-F-A-E-E-N-E-S-L-D-D-Q-N-I-S-I-A-G-H-T-F-I-D-R-P-N-Y-Q-F-T-N-L-K-A-A-K-K-G-S-M-V-Y-F-K-V-G-N-E-T-R-K-Y-K-M-T-S-I-R-D-V-K-P-T-D-V-G-V-L-D-E-Q-K-G-K-D-K-Q-L-T-L-I-T-C-D-D-Y-N-E-K-T-G-V-W-E-K-R-K-I-F-V-A-T-E-V-K (SEQ ID NO: 3) and (2) sequences incorporating one or more conservative amino acid substitutions in SEQ ID NO:3, wherein the conservative amino acid substitutions are any of the following: (1) any of isoleucine, leucine, and valine for any other of these amino acids; (2) aspartic acid for glutamic acid and vice versa; (3) glutamine for asparagine and vice versa; and (4) serine for threonine and vice versa.

Another aspect of the present invention is a nucleic acid sequence encoding this enzyme. In one alternative, the nucleic acid sequence includes therein a sequence of: (1)

ATGAAAAAATGGACAAATCGATTAATGACAATCGCTGGTGTGGTACTTATCCTAGTG
GCAGCATATTTGTTTGCTAAACCACATATCGATAATTATCTTCACGATAAAGATAAA
GATGAAAAGATTGAACAATATGATAAAAATGTAAAAGAACAGGCGAGTAAAGATA
AAAAGCAGCAAGCTAAACCTCAAATTCCGAAAGATAAATCGAAAGTGGCAGGCTAT
ATTGAAATTCCAGATGCTGATATTAAAGAACCAGTATATCCAGGACCAGCAACACCT
GAACAATTAAATAGAGGTGTAAGCTTTGCAGAAGAAAATGAATCACTAGATGATCA

AAATATTTCAATTGCAGGACACACTTTCATTGACCGTCCGAACTATCAATTTACAAA
TCTTAAAGCAGCCAAAAAAGGTAGTATGGTGTACTTTAAAGTTGGTAATGAAACAC
GTAAGTATAAAATGACAAGTATAAGAGATGTTAAGCCTACAGATGTAGGAGTTCTA
GATGAACAAAAAGGTAAAGATAAACAATTAACATTAATTACTTGTGATGATTACAA
TGAAAAGACAGGCGTTTGGGAAAAACGTAAAATCTTTGTAGCTACAGAAGTCAAAT
AA (SEQ ID NO: 2); and (2) a sequence complementary to SEQ ID NO: 2. In another alternative, the nucleic acid sequence can include a sequence hybridizing with SEQ ID NO: 2 or a sequence complementary to SEQ ID NO: 2 with no greater than about a 15% mismatch under stringent conditions. Preferably, the degree of mismatch is less than about 5%; more preferably, the degree of mismatch is less than about 2%.

Yet another aspect of the present invention is a vector comprising the nucleic acid sequence of the present invention operatively linked to at least one control sequence that controls the expression or regulation of the nucleic acid sequence.

Yet another aspect of the present invention is a host cell transfected with a vector of the present invention.

Another aspect of the present invention is a method for producing a substantially purified sortase-transamidase enzyme. The method comprises the steps of:

(1) culturing a host cell according to the present invention under conditions in which the host cell expresses the encoded sortase-transamidase enzyme; and

(2) purifying the expressed enzyme to produce substantially purified sortase-transamidase enzyme.

Another aspect of the present invention is a method for screening a compound for anti-sortase-transamidase activity. This method is important in providing a way to screen for antibiotics that disrupt the sorting reaction and are likely to be effective in treating infections caused by Gram-positive bacteria.

In one alternative, the screening method comprises the steps of:

(1) providing a substantially purified sortase-transamidase enzyme according to the present invention;

(2) performing an assay for sortase-transamidase in the presence and in the absence of the compound; and

(3) comparing the activity of the sortase-transamidase enzyme in the presence and in the absence of the compound to screen the compound for sortase-transamidase activity.

In another alternative, the screening method comprises the steps of:

(1) providing an active fraction of sortase-transamidase enzyme from a Gram-positive bacterium;

(2) performing an assay for sortase-transamidase in the presence and in the absence of the compound; and

(3) comparing the activity of the sortase-transamidase enzyme in the presence and in the absence of the compound to screen the compound for sortase-transamidase activity.

The active fraction of sortase-transamidase activity can be a particulate fraction from *Staphylococcus aureus* or another Gram-positive bacterium.

The assay for sortase-transamidase enzyme can be performed by monitoring the capture of a soluble peptide that is a substrate for the enzyme by its interaction with an affinity resin. In one alternative, the soluble peptide includes a sequence of at least six histidine residues and the affinity resin contains nickel. In another alternative, the soluble peptide includes the active site of glutathione S-transferase and the affinity resin contains glutathione. In yet another alternative, the soluble peptide includes the active site of streptavidin and the affinity resin contains biotin. In still another alternative, the soluble peptide includes the active site of maltose binding protein and the affinity resin contains amylose.

Still another aspect of the present invention is an antibody specifically binding the sortase-transamidase enzyme of the present invention.

Yet another aspect of the present invention is a protein molecule comprising a substantially purified sortase-transamidase enzyme according to the present invention extended at its carboxyl-terminus with a sufficient number of histidine residues to allow specific binding of the protein molecule to a nickel-sepharose column through the histidine residues added at the carboxyl-terminus.

Still another aspect of the present invention is a method for displaying a polypeptide on the surface of a Gram-positive bacterium comprising the steps of:

(1) expressing a polypeptide having a sorting signal at its carboxy-terminal end, the sorting signal having: (a) a motif of LPX_3X_4G therein; (b) a substantially hydrophobic domain of at least 31 amino acids carboxyl to the motif; and (c) a charged tail region with at least two positively charged residues carboxyl to the substantially hydrophobic domain, at least one of the two positively charged residues being arginine, the two positively charged residues being located at residues 31-33 from the motif, wherein X_3 is any of the twenty naturally-occurring L-amino acids and X_4 is selected from the group consisting of alanine, serine, and threonine;

(2) forming a reaction mixture including: (i) the expressed polypeptide; (ii) a substantially purified sortase-transamidase according to the present invention; and (iii) a Gram-positive bacterium having a peptidoglycan to which the sortase-transamidase can link the polypeptide; and

(3) allowing the sortase-transamidase to catalyze a reaction that cleaves the polypeptide within the LPX_3X_4G motif of the sorting signal and covalently cross-links the amino-terminal portion of the cleaved polypeptide to the peptidoglycan to display the polypeptide on the surface of the Gram-positive bacterium.

Another display method according to the present invention comprises:

(1) cloning a nucleic acid segment encoding a chimeric protein into a Gram-positive bacterium to generate a cloned chimeric protein including therein a carboxyl-terminal sorting signal as described above;

(2) growing the bacterium into which the nucleic acid segment has been cloned to express the cloned chimeric protein to generate a chimeric protein including therein a carboxyl-terminal sorting signal; and

(3) binding the polypeptide covalently to the cell wall by the enzymatic action of a sortase-transamidase expressed by the Gram-positive bacterium involving cleavage of the chimeric protein within the LPX_3X_4G motif so that the polypeptide is displayed on the surface of the Gram-positive bacterium in such a way that the polypeptide is accessible to a ligand.

Another aspect of the present invention is a polypeptide displayed on the surface of a Gram-positive bacterium by covalent linkage of an amino-acid sequence of LPX_3X_4 derived from cleavage of an LPX_3X_4G motif, wherein X_3 is any of the twenty naturally-occurring L-amino acids and X_4 is selected from the group consisting of alanine, serine, and threonine, the polypeptide being displayed on the surface of the Gram-positive bacterium in such a way that the polypeptide is accessible to a ligand.

Another aspect of the present invention is a covalent complex comprising:

(1) the displayed polypeptide; and

(2) an antigen or hapten covalently cross-linked to the polypeptide.

Yet another aspect of the present invention is a method for vaccination of an animal comprising the step of immunizing the animal with the displayed polypeptide to generate an immune response against the displayed polypeptide, or, alternatively, with the covalent complex to generate an immune response against the antigen or the hapten.

Still another aspect of the present invention is a method for screening for expression of a cloned polypeptide comprising the steps of:

(1) expressing a cloned polypeptide as a chimeric protein having a sorting signal at its carboxy-terminal end as described above;

(2) forming a reaction mixture including: (i) the expressed chimeric protein; (ii) a substantially purified sortase-transamidase enzyme according to the present invention; and (iii) a Gram-positive bacterium having a peptidoglycan to which the sortase-transamidase can link the polypeptide through the sorting signal;

(3) binding the chimeric protein covalently to the cell wall by the enzymatic action of a sortase-transamidase expressed by the Gram-positive bacterium involving cleavage of the chimeric protein within the LPX_3X_4G motif so that the polypeptide is displayed on the surface of the Gram-positive bacterium in such a way that the polypeptide is accessible to a ligand; and

(4) reacting the displayed polypeptide with a labeled specific binding partner to screen the chimeric protein for reactivity with the labeled specific binding partner.

Still another aspect of the present invention is a method for the diagnosis or treatment of a bacterial infection caused by a Gram-positive bacterium comprising the steps of:

(1) conjugating an antibiotic or a detection reagent to a protein including therein a carboxyl-terminal sorting signal as described above to produce a conjugate; and

(2) introducing the conjugate to an organism infected with a Gram-positive bacterium in order to cause the conjugate to be sorted and covalently cross-linked to the cell walls of the bacterium in order to treat or diagnose the infection.

If an antibiotic is used, typically it is a penicillin, ampicillin, vancomycin, gentamicin, streptomycin, a cephalosporin, amikacin, kanamycin, neomycin, paromomycin, tobramycin, ciprofloxacin, clindamycin, rifampin, chloramphenicol, norfloxacin, or a derivative of these antibiotics.

Similarly, another aspect of the present invention is a conjugate comprising an antibiotic or a detection reagent covalently conjugated to a protein including therein a carboxyl-terminal sorting signal as described above to produce a conjugate. In still another aspect of the present invention, a composition comprises the conjugate with a pharmaceutically acceptable carrier.

Another aspect of the present invention is a substantially purified protein having at least about 50% match with best alignment with the amino acid sequences of at least one of the putative homologous proteins of *Streptococcus pyogenes* (SEQ. ID NO. 4), *Actinomyces naeslundii* (SEQ. ID NO. 5), *Enterococcus faecalis* (SEQ. ID NO. 6), *Streptococcus mutans* (SEQ. ID. NO. 7) *or Bacillus subtilis* (SEQ. ID NO. 8) or *Streptococcus pneumoniae* (SEQ ID NO. ___) and having sortase-transamidase activity. Preferably, the match is at least about 60% in best alignment; more preferably, the match is at least about 70% in best alignment.

BRIEF DESCRIPTION OF THE DRAWINGS

These and other features, aspects, and advantages of the present invention will become better understood with reference to the following description and accompanying drawings where:

Figure 1 is a diagram of the activity of the sortase-transamidase enzyme of the present invention.

Figure 2:

(A) is a diagramatic representation of the primary structure of the surface protein precursor SEB-SPA490-524.

(B) depicts an SDS-PAGE gel of immunoprecipitated [^{35}S] SEB-SPA490-52 P1 precursor, P2 precursor and mature protein. SM317 and SM329 are two ts mutants that accumulate P2 as compared to wild-type staphylococci (WT).

(C) depicts an SDS-PAGE gel of immunoprecipitated [^{35}S] SEB-SPA490-52 P1 precursor, P2 precursor and mature protein in SM317, SM329 and WT staphylococci following a pulse-chase analysis of SEB-SPA490-524 anchoring.

(D) depicts Staphylococcal strains OS2 (WT), SM317 and SM329 streaked on tryptic soy agar and grown at 42°C.

Figure 3:

(A) is a diagrammatic representation of the primary structure of SEB-MH$_6$-CWS and its linkage to the cell wall.

(B) deptics a mass spectroscopy profile (MALDI-MS) of solubilized and affinity purified SEB-MH$_6$-CWS.

(**C**) deptics a mass spectroscopy profile (MALDI-MS) of solubilized, mutanolysin-released anchor peptides were digested with f11 hydrolase.

Figure 4:

(**A**) depicts an SDS-PAGE gel of immunoprecipitated [^{35}S] SEB-SPA490-52 P1 precursor, P2 precursor and mature protein in SM317, SM329 and WT staphylococci transformed with or without pGL1834 (plasmid containing the *srtA* gene cloned into pC194-mcs) following a pulse-chase analysis of SEB-SPA490-524 anchoring.

(**B**) depicts an SDS-PAGE gel of immunoprecipitated [^{35}S] SEB-SPA490-52 P1 precursor, P2 precursor and mature protein from SM317 transformed with the DNA of either the mutant SM317 (pGL1898) or wild-type strain OS2 (pGL1897).

(**C**) depicts an SDS-PAGE gel of immunoprecipitated [35S] SEB-SPA490-52 P1 precursor, P2 precursor and mature protein from *S. aureus* OS2 (wild type), SM317 and SM329 transformed with pGL1834 and subjected to pulse-chase analysis.

Figure 5 depicts the size of DNA fragments and the position of the coding region of the *srtA* gene of *S. aureus* (SEQ ID NO: 2) sufficient for an increase in surface protein anchoring. The concentration of P2 precursor in plasmid transformants of the mutant SM317 was measured by labeling with [^{35}S]methionine and is indicated in percent.

Figure 6 depicts the DNA sequence of the *srtA* gene (SEQ ID NO: 2) and deduced primary structure of the SrtA protein (SEQ ID NO: 3). The NH_2-terminal hydrophobic membrane anchor sequence is boxed. A single cysteine predicted to be the active site for cleavage of cell wall sorting signals at the LPXTG motif is shaded.

Figure 7 depicts a sequence alignment comparing the predicted primary structure of the SrtA protein (Sortase) with that of homologous sequences identified by database searches. Note the conservation of a single cysteine residue as well as its surrounding sequence.

Figure 8:

(A) depicts the structure of Seb-Spa490-524 harboring an NH_2-terminal leader (signal) peptide with signal peptidase cleavage site as well as a COOH-terminally fused cell wall sorting signal consisting of the LPXTG motif, hydrophobic domain (*black box*), and positively charged tail (*boxed* +).

(B) depicts the SDS-PAGE gel analysis of pulse chase experiment where staphlococcal cultures were labeled with [^{35}S]methionine for 1 min and quenching all further incorporation by the addition of excess unlabeled methionine (chase). P1 precursor, P2 precursor and mature Seb-Spa490-524 were evaluated.

Figure 9:

(A) depicts a growth curve for staphylococcal growth with antibiotics added (1, open squares: mock treated; 2, open diamonds: penicillin 10 μg/ml; 3, closed diamonds: moenomycin, 10 μg/ml; 4, closed squares: vancomycin 10 μg/ml).

(B) depicts a curve measuring the rate of cell wall sorting in the presence of antibiotics or mock treated as described in **(A)**.

Figure 10:

(A) depicts the structure of Seb-Cws-BlaZ harboring an NH_2-terminal signal (leader) peptide and the sorting signal of protein A which consists of an LPXTG motif, hydrophobic (*shaded box*) and charged domains (*boxed* RRREL). The sorting signal is fused to the COOH-terminus of Seb and to the NH_2-terminus of mature BlaZ. Cleavage at the LPXTG motif produces two fragments, an NH_2-terminal cell wall anchored surface protein (Seb) and a COOH-terminal BlaZ domain that is located in the bacterial cytoplasm.

(B) depicts an SDS-PAGE gel analysis of *S. aureus* OS2 (pSeb-Cws-BlaZ) and *S. aureus* OS2 (pSeb-Cws$_{DLPXTG}$-BlaZ) cell wall sorting. The arrows point to Seb species that were observed in protoplasts but not in whole cells.

Figure 11 depicts a model for the transpeptidation reaction catalyzed by staphylococcal sortase.

Figure 12:

(A) depicts an SDS-PAGE gel analysis of a pulse chance analysis of surface protein anchoring to the cell wall in the presence or absence of release of proteins fro the surface by hydroxylamine.

(B) depicts an SDS-PAGE gel analysis of a pulse chance analysis of surface protein anchoring to the cell wall in the presence or absence of release of proteins fro the surface by hydroxylamine added either 5 min prior to labeling (prior), during pulse-labeling (pulse) or 5 min after quenching to *S. aureus* OS2 cultures.

(**C**) depicts a bar graph indicating that increasing amounts of hydroxylamine added 5 min prior to labeling of *S. aureus* OS2 cultures caused increasing amounts of surface protein to be released.

Figure 13:

(**A**) depicts a Coomassie-stained SDS-PAGE gel used to characterize surface proteins released by hydroylamine treatment.

(**B**) depicts an rpHPLC chromatogram of COOH-terminal anchor peptides released from *S. aureus* BB270 cells via treatment with 0.1 M NH_2OH.

(**C**) depicts an rpHPLC chromatogram of COOH-terminal anchor peptides released from *S. aureus* BB270 cells via treatment with 0.1 M NH_2OH.

Figure 14:

(**A**) is a bar graph depicting the effect of incubating staphyloccocal extracts with the sorting substrate DABCYL-QALPETGEE-EDANS; peptide cleavage is indicated as an increase in fluorescence. The addition of 0.2 M NH_2OH increased peptide cleavage, whereas peptide cleavage was inhibited by the addition of methanethiosulfonate (MTSET), a known inhibitor of sortase.

(**B**) depicts an SDS-PAGE gel analysis of *E. coli* XL-1Blue (pHTT5) expressing $SrtA_{DN}$, in which the NH_2-terminal membrane anchor of sortase (SrtA) has been replaced with a six histidine tag. Lane 1 contains uninduced culture; 2, 1 mM IPTG induced culture; 3, French press extract; 4, the supernatant of centrifuged French press extracts; 5, the sediment of French press extracts; 6, flow-through of affinity chromatography on Ni-NTA; 7, column wash; 8-10, 1 ml fractions eluted with 0.5 M imidazol.

(C) is a bar graph depicting the effect of incubating purified $SrtA_{DN}$ with the peptide substrate DABCYL-QALPETGEE-EDANS and cleavage monitored as an increase in fluorescence. The reaction was inhibited by the addition of methanethiosulfonate (MTSET) or organic mercurial (pHMB), while the addition of 0.2 M NH_2OH accelerated cleavage. MTSET-treated $SrtA_{DN}$ could be rescued by incubation with 10 mM DTT.

DEFINITIONS

As used herein, the terms defined below have the following meanings unless otherwise indicated:

"Nucleic Acid Sequence": the term "nucleic acid sequence" includes both DNA and RNA unless otherwise specified, and, unless otherwise specified, includes both double-stranded and single-stranded nucleic acids. Also included are hybrids such as DNA-RNA hybrids. In particular, a reference to DNA includes RNA that has either the equivalent base sequence except for the substitution of uracil and RNA for thymine in DNA, or has a complementary base sequence except for the substitution of uracil for thymine, complementarity being determined according to the Watson-Crick base pairing rules. Reference to nucleic acid sequences can also include modified bases as long as the modifications do not significantly interfere either with binding of a ligand such as a protein by the nucleic acid or with Watson-Crick base pairing.

"Mismatch": as used herein the term "mismatch" includes all unpaired bases when two nucleic acid sequences are hybridized with best alignment in the context of nucleic acid hybridization. In other words, the term "mismatch" includes not only situations in which the same number of bases are present in the two sequences or segments of sequences, but in which some bases do not form Watson-Crick pairs because of their sequences, but also situations in which different numbers of bases are present in the two sequences because of insertions or deletions, referred to generically as "indels." In this latter situation, certain of the bases in the longer sequence must be unpaired and may loop out from the hybrid.

"Match": as used herein the term "match" includes all paired amino acids when two amino acid sequences are compared with best alignment in the context in terms of protein sequence comparison. Amino acid "sequence identity" percentages include only identical amino acid pairing when amino acid sequences are matched in best alignment. Amino acid "sequence similarity" percentages include both similar and identical amino acids when amino acid sequences are matched in best alignment. Similar amino acids are amino acids which share similar physical and/or chemical properties. The following is a listing of amino acids which are

considered to be similar, or conservative amino acids relative to one another, as substitutions of each of these amino acids for the other in a sequence often do not disrupt the structure or function of the molecule as the amino acids share similar physical and/or chemical properties. In particular, the conservative amino acid substitutions can be any of the following: (1) any of isoleucine for leucine or valine, leucine for isoleucine, and valine for leucine or isoleucine; (2) aspartic acid for glutamic acid and glutamic acid for aspartic acid; (3) glutamine for asparagine and asparagine for glutamine; and (4) serine for threonine and threonine for serine.

Other substitutions can also be considered conservative, depending upon the environment of the particular amino acid. For example, glycine (G) and alanine (A) can frequently be interchangeable, as can be alanine and valine (V). Methionine (M), which is relatively hydrophobic, can frequently be interchanged with leucine and isoleucine, and sometimes with valine. Lysine (K) and arginine (R) are frequently interchangeable in locations in which the significant feature of the amino acid residue is its charge and the different pK's of these two amino acid residues or their different sizes are not significant. Still other changes can be considered "conservative" in particular environments. For example, if an amino acid on the surface of a protein is not involved in a hydrogen bond or salt bridge interaction with another molecule, such as another protein subunit or a ligand bound by the protein, negatively charged amino acids such as glutamic acid and aspartic acid can be substituted for by positively charged amino acids such as lysine or arginine and vice versa. Histidine (H), which is more weakly basic than arginine or lysine, and is partially charged at neutral pH, can sometimes be substituted for these more basic amino acids. Additionally, the amides glutamine (Q) and asparagine (N) can sometimes be substituted for their carboxylic acid homologues, glutamic acid and aspartic acid.

"Antibody": as used herein the term "antibody" includes both intact antibody molecules of the appropriate specificity, and antibody fragments (including Fab, F(ab'), Fv, and $F(ab')_2$), as well as chemically modified intact antibody molecules and antibody fragments, including hybrid antibodies assembled by in vitro reassociation of subunits. Also included are single-chain antibody molecules generally denoted by the term sFv and humanized antibodies in which some or all of the originally non-human constant regions are replaced with constant

regions originally derived from human antibody sequences. Both polyclonal and monoclonal antibodies are included unless otherwise specified. Additionally included are modified antibodies or antibodies conjugated to labels or other molecules that do not block or alter the binding capacity of the antibody.

DESCRIPTION

A substantially purified sortase-transamidase enzyme from Gram-positive bacteria, particularly *Staphylococcus aureus*, has been identified and purified. The properties of this enzyme make it a logical target for antibiotic action. This enzyme also catalyzes covalent crosslinkage of proteins to the peptidoglycan of Gram-positive bacteria.

I. THE SORTASE-TRANSAMIDASE ENZYME

One aspect of the invention is a substantially purified sortase-transamidase enzyme from a Gram-positive bacterium. As used herein, the term "substantially purified" means having a specific activity of at least tenfold greater than the sortase-transamidase activity present in a crude extract, lysate, or other state from which proteins have not been removed and also in substantial isolation from proteins found in association with sortase-transamidase in the cell.

The enzyme has a molecular weight of about 23,539 daltons. The enzyme catalyzes a reaction that covalently crosslinks the carboxyl-terminus of a protein having a sorting signal to the peptidoglycan of the Gram-positive bacterium. The sorting signal has: (1) a motif of LPX_3X_4G therein; (2) a substantially hydrophobic domain of at least 31 amino acids carboxyl to the motif; and (3) a charged tail region with at least two positively charged residues carboxyl to the substantially hydrophobic domain, at least one of the two positively charged residues being arginine, the two positively charged residues being located at residues 31-33 from the motif. In this sorting signal, X_3 can be any of the twenty naturally-occurring L-amino acids. X_4 can be alanine, serine, or threonine. Preferably, X_4 is threonine.

The sortase-transamidase is believed to occur in all Gram-positive bacteria. In particular, the enzyme exists in *Mycobacterium*, *Nocardia*, *Actinomyces*, *Staphylococcus*, *Streptococcus*, *Listeria*, *Enterococcus*, *Bacillus*, and *Pneumococcus*. Specifically, the enzyme exists in the following species: *Staphylococcus aureus*, *S. sobrinus*, *Enterococcus faecalis*, *Streptococcus pyogenes*, *Bacillus subtilis*, *Streptococcus pneumoniae,* and *Listeria monocytogenes*.

Preferably the enzyme is isolated from *Staphylococcus aureus*, and more preferably is a product of the *srt*A gene of *S. aureus*.

A. Amino Acid Sequence

The sortase-transamidase of the present invention includes therein an amino acid sequence of: M-K-K-W-T-N-R-L-M-T-I-A-G-V-V-L-I-L-V-A-A-Y-L-F-A-K-P-H-I-D-N-Y-L-H-D-K-D-K-D-E-K-I-E-Q-Y-D-K-N-V-K-E-Q-A-S-K-D-K-K-Q-Q-A-K-P-Q-I-P-K-D-K-S-K-V-A-G-Y-I-E-I-P-D-A-D-I-K-E-P-V-Y-P-G-P-A-T-P-E-Q-L-N-R-G-V-S-F-A-E-E-N-E-S-L-D-D-Q-N-I-S-I-A-G-H-T-F-I-D-R-P-N-Y-Q-F-T-N-L-K-A-A-K-K-G-S-M-V-Y-F-K-V-G-N-E-T-R-K-Y-K-M-T-S-I-R-D-V-K-P-T-D-V-G-V-L-D-E-Q-K-G-K-D-K-Q-L-T-L-I-T-C-D-D-Y-N-E-K-T-G-V-W-E-K-R-K-I-F-V-A-T-E-V-K (SEQ ID NO: 3).

Also within the scope of the present invention are substantially purified protein molecules that are mutants of the sequence of SEQ ID NO:3 that preserve the sortase-transamidase activity. In particular, the conservative amino acid substitutions can be any of the following: (1) any of isoleucine for leucine or valine, leucine for isoleucine, and valine for leucine or isoleucine; (2) aspartic acid for glutamic acid and glutamic acid for aspartic acid; (3) glutamine for asparagine and asparagine for glutamine; and (4) serine for threonine and threonine for serine.

Other substitutions can also be considered conservative, depending upon the environment of the particular amino acid. For example, glycine (G) and alanine (A) can frequently be interchangeable, as can be alanine and valine (V). Methionine (M), which is relatively hydrophobic, can frequently be interchanged with leucine and isoleucine, and

sometimes with valine. Lysine (K) and arginine (R) are frequently interchangeable in locations in which the significant feature of the amino acid residue is its charge and the different pK's of these two amino acid residues or their different sizes are not significant. Still other changes can be considered "conservative" in particular environments. For example, if an amino acid on the surface of a protein is not involved in a hydrogen bond or salt bridge interaction with another molecule, such as another protein subunit or a ligand bound by the protein, negatively charged amino acids such as glutamic acid and aspartic acid can be substituted for by positively charged amino acids such as lysine or arginine and vice versa. Histidine (H), which is more weakly basic than arginine or lysine, and is partially charged at neutral pH, can sometimes be substituted for these more basic amino acids. Additionally, the amides glutamine (Q) and asparagine (N) can sometimes be substituted for their carboxylic acid homologues, glutamic acid and aspartic acid.

The amino acid sequence (SEQ ID NO: 3) of sortase-transamidase from *Staphylococcus aureus* has substantial homology with sequences of enzymes from other Gram-positive bacteria. There is about a 31% sequence identity (and about 44% sequence simlarity) with best alignment over the entire sequenced region of the *S. pyogenes* open reading frame (SEQ. ID NO. 4). There is about a 28% sequence identity (and about 44% sequence similarity) with best alignment over the entire sequenced region of the *A. naeslundii* open reading frame (SEQ. ID NO. 5). There is about a 27% sequence identity (and about 47% sequence similarity) with best alignment over the entire sequenced region of the *S. mutans* open reading frame (SEQ. ID NO. 7). There is about a 25% sequence identity (and about 45% sequence similarity) with best alignment over the entire sequenced region of the *E. faecalis* open reading frame (SEQ. ID NO. 6). There is about a 23% identity and about a 38% similarity between the sequence with best alignment over the entire sequenced region of the *B. subtilis* open reading frame (SEQ. ID NO. 8). as compared with the *S.* mutans open reading frame (SEQ. ID NO. 7), with a lower degree of sequence identity and similarity between the *B. subtilis* and *S. pyogenes* open reading frames. These matches are shown in Figure 7.

Additionally, there is about a 32% sequence identity and about a 47% sequence similarity with best alignment over the entire sequence between the *S. aureus* open reading frame

(SEQ. ID NO. 3) and a protein designated srtA from *Streptococcus pneumoniae* (SEQ. ID NO. 34). The sequence of the srtA protein is

MSRTKLRALLGYLLMLVACLIPIYCFGQMVLQSLGQVKGHATFVKSMTTEMYQEQQN
HSLAYNQRLASQNRIVDPFLAEGYEVNYQVSDDPDAVYGYLSIPSLEIMEPVYLGADYH
HLGMGLAHVDGTPLPLDGTGIRSVIAGHRAEPSHVFFRHLDQLKVGDALYYDNGQEIVE
YQMMDTEIILPSEWEKLESVSSKNIMTLITCDPIPTFNKRLLVNFERVAVYQKSDPQTAA
VARVAFTKEGQSVSRVATSQWLYRGLVVLAFLGILFVLWKLARLLRGK (SEQ ID NO. 34). Similarly, there is about a 30% sequence identity and about a 46% sequence similarity with best alignment over the entire sequence between the *S. aureus* open reading frame (SEQ. ID NO.3) and a protein designated srtB from *Streptococcus pneumoniae* (SEQ. ID NO. 35). The sequence of the srtB protein is

MDNSRRSRKKGTKKKKHPLILLLIFLVGFAVAIYPLVSRYYYRISNEVIKEFDETVSQMD
KAELEERWRLAQAFNATLKPSEILDPFTEQEKKKGVSEYANMLKVHERIGYVEIPAIDQE
IPMYVGTSEDILQKGAGLLEGASLPVGGENTHTVITAHRGLPTAELFSQLDKMKKGDIFY
LHVLDQVLAYQVDQIVTVEPNDFEPVLIQHGEDYATLLTCTPYMINSHRLLVRGKRIPYT
APIAERNRAVRERGQFWLWLLLGAMAVILLLLYRVYRNRRIVKGLEKQLEGRHVKD (SEQ. ID NO. 35). Similarly, there is about a 29% sequence identity and about a 43% sequence similarity with best alignment over the entire sequence between the *S. aureus* open reading frame (SEQ. ID NO.3) and a protein designated srtC from *Streptococcus pneumoniae* (SEQ ID NO. 36). The sequence of the srtC protein is

MDNSRRSRKKGTKKKKHPLILLLIFLVGFAVAIYPLVSRYYYRISNEVIKEFDETVSQMD
KAELEERWRLAQAFNATLKPSEILDPFTEQEKKKGVSEYANMLKVHERIGYVEIPAIDQE
IPMYVGTSEDILQKGAGLLEGASLPVGGENTHTVITAHRGLPTAELFSQLDKMKKGDIFY
LHVLDQVLAYQVDQIVTVEPNDFEPVLIQHGEDYATLLTCTPYMINSHRLLVRGKRIPYT
APIAERNRAVRERGQFWLWLLLGAMAVILLLLYRVYRNRRIVKGLEKQLEGRHVKD (SEQ ID NO. 36).

Therefore, another aspect of the present invention is a substantially purified protein molecule that has at least a 18% sequence identity match, preferably a 20% sequence

identity match, and most preferably a 30% sequence identity match with best alignment with the *S. pyogenes, A. naeslundii, S. mutans, E. faecalis* or *B. subtilis* open reading frame of Figure 7 and that has sortase-transamidase activity. Further, another aspect of the present invention is a substantially purified protein molecule that has at least a 30% sequence similarity match, preferably a 40% sequence similarity match, and most preferably a 50% sequence similarity match with best alignment with the *S. pyogenes, A. naeslundii, S. mutans, E. faecalis* or *B. subtilis* open reading frame of Figure 7 and that has sortase-transamidase activity.

The sortase-transamidase is a cysteine protease.

B. Activity of the Sortase-Transamidase

The activity of the sortase-transamidase enzyme of the present invention is shown, in general, in Figure 1. The enzyme first cleaves a polypeptide having a sorting signal within the LPX_3X_4G motif. Cleavage occurs after residue X_4, normally a threonine; as indicated above, this residue can also be a serine or alanine residue. This residue forms a covalent intermediate with the sortase. The next step is the transamidation reaction that transfers the cleaved carboxyl terminus of the protein to be sorted to the $-NH_2$ of the pentaglycine crossbridge within the peptidoglycan precursor. The peptidoglycan precursor is then incorporated into the cell wall by a transglycosylase reaction with the release of undecaprenyl phosphate. The mature anchored polypeptide chains are thus linked to the pentaglycine cross bridge in the cell wall which is tethered to the ε-amino side chain of an unsubstituted cell wall tetrapeptide. A carboxypeptidase may cleave a D-Ala-D-Ala bond of the pentapeptide structure to yield the final branched anchor peptide in the staphylococcal cell wall.

The sorting signal has: (1) a motif of LPX_3X_4G therein; (2) a substantially hydrophobic domain of at least 31 amino acids carboxyl to the motif; and (3) a charged tail region.

In the motif, X_3 can be any of the 20 naturally-occurring L-amino acids. X_4 can be any of threonine, serine, or alanine. Preferably, X_4 is threonine (O. Schneewind et al., "Cell

Wall Sorting Signals in Surface Proteins of Gram-Positive Bacteria," EMBO J. 12:4803-4811 (1993)).

Preferably, the substantially hydrophobic domain carboxyl to the motif includes no more than about 7 charged residues or residues with polar side chains. For the purposes of this specification, these residues include the following: aspartic acid, glutamic acid, lysine, and arginine as charged residues, and serine, threonine, glutamine, and asparagine as polar but uncharged residues. Preferably, the sequence includes no more than three charged residues.

Representative sequences suitable for sorting signals for use with the sortase-transamidase of the present invention include, but are not limited to the following: E-E-N-P-F-I-G-T-T-V-F-G-G-L-S-L-A-L-G-A-A-L-L-A-G (SEQ ID NO: 9), the hydrophobic domain of the staphylococcal proteinase (SPA) sorting signal from *Staphylococcus aureus*; (2) G-E-E-S-T-N-K-G-M-L-F-G-G-L-F-S-I-L-G-L-A-L-L (SEQ ID NO:10), the SNBP signal of *S. aureus*; (3) D-S-S-N-A-Y-L-P-L-L-G-L-V-S-L-T-A-G-F-S-L-L-G-L (SEQ ID NO: 11), the SPAA signal of *S. sobrinus*, (4) E-K-Q-N-V-L-L-T-V-V-G-S-L-A-A-M-L-G-L-A-G-L-G-F (SEQ ID NO:12), the PRGB signal of *Enterococcus faecalis*, (5) S-I-G-T-Y-L-F-K-I-G-S-A-A-M-I-G-A-I-G-I-Y-I-V (SEQ ID NO:13), the TEE signal of *Streptococcus pyogenes*, and (6) D-S-D-N-A-L-Y-L-L-L-G-L-L-A-V-G-T-A-M-A-L-T (SEQ ID NO:14), the INLA signal of *Listeria monocytogenes*. Other hydrophobic domains can be used as part of the sorting signal.

The third portion of the sorting signal is a charged tail region with at least two positively charged residues carboxyl to the substantially hydrophobic domain. At least one of the two positively charged residues is arginine. The charged tail can also contain other charged amino acids, such as lysine. Preferably, the charged tail region includes two or more arginine residues. The two positively charged residues are located at residues 31-33 from the motif. Preferably, the two arginine residues are either in succession or are separated by no more than one intervening amino acid. Preferably, the charged tail is at least five amino acids long, although four is possible. Among the charged tails that can be used are the following: (1) R-R-R-E-L (SEQ ID NO:15), from the SPA signal of *S. aureus*; (2) R-R-N-K-K-N-H-K-A (SEQ ID NO:16), from the SNBP signal of *S. aureus*; (3) R-R-K-Q-D (SEQ ID NO:17), from the SPAA

signal of *S. sobrinus*; (4) K-R-R-K-E-T-K (SEQ ID NO:18), from the PRGB signal of *E. faecalis*; (5) K-R-R-K-A (SEQ ID NO:19), from the TEE signal of *S. pyogenes*; (6), K-R-R-H-V-A-K-H (SEQ ID NO:20), from the FIM sorting signal of *Actinomyces viscosus*, and (7) K-R-R-K-S (SEQ ID NO:21), from the BAC sorting signal of *Streptococcus aglactiae*; (8) K-R-K-E-E-N (SEQ ID NO:22), from the EMM signal of *Streptococcus pyogenes*.

Also usable as the charged tail portion of the sorting signal are the following sequences produced by mutagenesis from the SPA signal of *S. aureus*. These include R-R-R-E-S (SEQ ID NO: 23), R-R-R-S-L (SEQ ID NO: 24), R-R-S-E-L (SEQ ID NO: 25), R-S-R-E-L (SEQ ID NO: 26) and S-R-R-E-L (SEQ ID NO: 27). Other charged tails that are usable as part of the sorting signal can be derived from a polyserine tail, itself inactive, by replacement of one or more of the serine residues with the basic amino acid arginine. These include R-R-S-S-S (SEQ ID NO: 28), R-S-R-S-S (SEQ ID NO:29), and S-R-R-S-S (SEQ ID NO:30). Other sorting signals can also be used.

II. THE GENE ENCODING THE SORTASE-TRANSAMIDASE ENZYME

A. Isolation of the Sortase-Transamidase Enzyme Gene

The gene for the sortase-transamidase enzyme in *Staphylococcus aureus*, the *srt*A gene, has been isolated. The isolation process is described in detail in the Example below; in general, this process comprises: (1) the generation of temperature-sensitive mutants through chemical mutagenesis, such as with the DNA modifying agent N-methyl-N-nitro-N-nitrosoguanidine; (2) screening for temperature-sensitive mutants; (3) screening the temperature-sensitive mutants for a block in protein sorting by the use of a construct harboring the staphylococcal enterotoxin B (SEB) gene fused to the cell wall sorting signal of staphylococcal Protein A (SPA), to locate mutants that accumulate a precursor molecule formed by cleavage of an amino-terminal signal peptide but that is not then processed by cleavage of the carboxyl-terminal sorting signal; (4) generation of a *S. aureus* chromosomal library and complementation of the sorting defect leading to abnormal accumulation of the P2 precursor; and (5) sequencing and characterization of the *S. aureus* complementing determinants.

B. Sequence of the Sortase-Transamidase Gene

The above procedure yielded the entire coding sequence for the sortase-transamidase gene, *srt*A. This sequence is:

ATGAAAAAATGGACAAATCGATTAATGACAATCGCTGGTGTGGTACTTATCCTAGTG
GCAGCATATTTGTTTGCTAAACCACATATCGATAATTATCTTCACGATAAAGATAAA
GATGAAAAGATTGAACAATATGATAAAAATGTAAAAGAACAGGCGAGTAAAGATA
AAAAGCAGCAAGCTAAACCTCAAATTCCGAAAGATAAATCGAAAGTGGCAGGCTAT
ATTGAAATTCCAGATGCTGATATTAAAGAACCAGTATATCCAGGACCAGCAACACCT
GAACAATTAAATAGAGGTGTAAGCTTTGCAGAAGAAAATGAATCACTAGATGATCA
AAATATTTCAATTGCAGGACACACTTTCATTGACCGTCCGAACTATCAATTTACAAA
TCTTAAAGCAGCCAAAAAAGGTAGTATGGTGTACTTTAAAGTTGGTAATGAAACAC
GTAAGTATAAAATGACAAGTATAAGAGATGTTAAGCCTACAGATGTAGGAGTTCTA
GATGAACAAAAAGGTAAAGATAAACAATTAACATTAATTACTTGTGATGATTACAA
TGAAAAGACAGGCGTTTGGGAAAAACGTAAAATCTTTGTAGCTACAGAAGTCAAAT
AA (SEQ ID NO: 2). The last three nucleotides, TAA, of this sequence are the stop codon.

Accordingly, within the scope of the present invention is a nucleic acid sequence encoding a substantially purified sortase-transamidase enzyme from a Gram-positive bacterium. The enzyme encoded has a molecular weight of about 23,539 daltons and catalyzes a reaction that covalently cross-links the carboxyl-terminus of a protein having the sorting signal described above to the peptidoglycan of a gram-positive bacterium. The nucleic acid sequence includes therein the sequence of SEQ ID NO: 2 or a sequence complementary to SEQ ID NO: 2.

Also included within the present invention is a nucleic acid sequence encoding a substantially purified sortase-transamidase enzyme from a Gram-positive bacterium with a molecular weight of about 23,539 daltons, where the enzyme catalyzes the cross-linking reaction where the nucleic acid sequence hybridizes with at least one of: (1) the sequence of SEQ ID NO: 2; (2) a sequence complementary to SEQ ID NO: 2; or (3) a sequence complementary to SEQ ID NO: 2 with no greater than about a 15% mismatch under stringent conditions. Preferably, the

degree of mismatch is no greater than about 5%; most preferably the mismatch is no greater than about 2%.

Also within the present invention is a nucleic acid sequence encoding a substantially purified sortase-transamidase enzyme from a Gram-positive bacterium with a molecular weight of about 23,539 daltons and catalyzes the cross-linking reaction described above involving the sorting signal, where the enzyme includes therein an amino acid sequence selected from the group consisting of: (1) M-K-K-W-T-N-R-L-M-T-I-A-G-V-V-L-I-L-V-A-A-Y-L-F-A-K-P-H-I-D-N-Y-L-H-D-K-D-K-D-E-K-I-E-Q-Y-D-K-N-V-K-E-Q-A-S-K-D-K-K-Q-Q-A-K-P-Q-I-P-K-D-K-S-K-V-A-G-Y-I-E-I-P-D-A-D-I-K-E-P-V-Y-P-G-P-A-T-P-E-Q-L-N-R-G-V-S-F-A-E-E-N-E-S-L-D-D-Q-N-I-S-I-A-G-H-T-F-I-D-R-P-N-Y-Q-F-T-N-L-K-A-A-K-K-G-S-M-V-Y-F-K-V-G-N-E-T-R-K-Y-K-M-T-S-I-R-D-V-K-P-T-D-V-G-V-L-D-E-Q-K-G-K-D-K-Q-L-T-L-I-T-C-D-D-Y-N-E-K-T-G-V-W-E-K-R-K-I-F-V-A-T-E-V-K (SEQ ID NO: 3); and (2) sequences incorporating one or more conservative amino acid substitutions in SEQ ID NO:3 wherein the conservative amino acid substitutions are any of the following: (1) any of isoleucine, leucine and valine for any other of these amino acids; (2) aspartic acid for glutamic acid and vice versa; (3) glutamine for asparagine and vice versa; and (4) serine for threonine and vice versa. Alternative nucleic acid sequences can be determined using the standard genetic code; the alternative codons are readily determinable for each amino acid in this sequence.

Construction of nucleic acid sequences according to the present invention can be accomplished by techniques well known in the art, including solid-phase nucleotide synthesis, the polymerase chain reaction (PCR) technique, reverse transcription of DNA from RNA, the use of DNA polymerases and ligases, and other techniques. If an amino acid sequence is known, the corresponding nucleic acid sequence can be constructed according to the genetic code.

C. Vectors and Host Cells Transformed with Vectors

Another aspect of the invention is a vector comprising a nucleic acid sequence according to the present invention operatively linked to at least one control sequence that controls the expression or regulation of the nucleic acid sequence. Such control sequences are

well known in the art and include operators, promoters, enhancers, promoter-proximal elements and replication origins. The techniques of vector construction, including cloning, ligation, gap-filling, the use of the polymerase chain reaction (PCR) procedure, solid-state oligonucleotide synthesis, and other techniques, are all well known in the art and need not be described further here.

Another aspect of the present invention is a host cell transfected with a vector according to the present invention. Among the host cells that can be used are gram-positive bacteria such as *Staphylococcus aureus*.

Transfection, also known as transformation, is done using standard techniques appropriate to the host cell used, particularly *Staphylococcus aureus*. Such techniques are described, for example, in R.P. Novick, "Genetic Systems in Staphylococci," Meth. Enzymol. 204: 587-636 (1991), as well as in O. Schneewind et al., "Sorting of Protein A to the Staphylococcal Cell Wall," Cell 70: 267-281 (1992).

III. SORTASE-TRANSAMIDASE AS A TARGET FOR ANTIBIOTIC ACTION

A. A Site for Antibiotic Action

The reaction carried out by the sortase-transamidase of the present invention presents a possible target for a new class of antibiotics to combat medically relevant infections caused by numerous gram-positive organisms. Because this is a novel site of antibiotic action, these antibiotics have the advantage that resistance by the bacterium has not had a chance to develop.

Such antibiotics can include compounds with structures that mimic the cleavage site, such as compounds with a structure similar to methyl methanethiosulfonate or, more generally, alkyl methanethiosulfonates. The sortase-transamidase of the present invention is believed to be a cysteine protease. Other antibiotics that may inhibit the activity of the sortase-transamidase in the present invention include inhibitors that would be specific for cysteine-modification in a β-lactam framework. These inhibitors can, but need not necessarily, have active moieties that would form mixed disulfides with the cysteine sulfhydryl. These active

moieties could be derivatives of methanethiosulfonate, such as methanethiosulfonate ethylammonium, methanethiosulfonate ethyltrimethylammonium, or methanethiosulfonate ethylsulfonate (J.A. Javitch et al., "Mapping the Binding Site Crevice of the Dopamine D2 Receptor by the Substituted-Cysteine Accessibility Method," Neuron, 14: 825-831 (1995); M.H. Akabas & A. Karlin, "Identification of Acetylcholine Receptor Channel-Lining Residues in the M1 Segment of the α-Subunit," Biochemistry 34: 12496-12500 (1995)). Similar reagents, such as alkyl alkanethiosulfonates, i.e., methyl methanethiosulfonate, or alkoxycarbonylalkyl disulfides, have been described (D.J. Smith et al., "Simple Alkanethiol Groups for Temporary Blocking of Sulfhydryl Groups of Enzymes," Biochemistry 14: 766-771 (1975); W.N. Valentine & D.E. Paglia, "Effect of Chemical Modification of Sulfhydryl Groups of Human Erythrocyte Enzymes," Am. J. Hematol. 11: 111-124 (1981)). Other useful inhibitors involve derivatives of 2-trifluoroacetylaminobenzene sulfonyl fluoride (J.C. Powers, "Proteolytic Enzymes and Their Active-Site-Specific Inhibitors: Role in the Treatment of Disease," in Modification of Proteins), in a β-lactam framework, peptidyl aldehydes and nitriles (E. Dufour et al., "Peptide Aldehydes and Nitriles as Transition State Analog Inhibitors of Cysteine Proteases," Biochemistry 34: 9136-9143 (1995); J. O. Westerik & R. Wolfenden, "Aldehydes as Inhibitors of Papain," J. Biol. Chem. 247: 8195-8197 (1972)), peptidyl diazomethyl ketones (L. Björck et al., "Bacterial Growth Blocked by a Synthetic Peptide Based on the Structure of a Human Proteinase Inhibitor," Nature 337: 385-386 (1989)), peptidyl phosphonamidates (P.A. Bartlett & C.K. Marlowe, "Phosphonamidates as Transition-State Analogue Inhibitors of Thermolysin," Biochemistry 22: 4618-4624 (1983)), phosphonate monoesters such as derivatives or analogues of m-carboxyphenyl phenylacetamidomethylphosphonate (R.F. Pratt, "Inhibition of a Class C β-Lactamase by a Specific Phosphonate Monoester," Science 246: 917-919 (1989)), maleimides and their derivatives, including derivatives of such bifunctional maleimides as o-phenylenebismaleimide, p-phenylenebismaleimide, m-phenylenebismaleimide, 2,3-naphthalenebismaleimide, 1,5-naphthalenebismaleimide, and azophenylbismaleimide, as well as monofunctional maleimides and their derivatives (J.V. Moroney et al., "The Distance Between Thiol Groups in the γ Subunit of Coupling Factor 1 Influences the Proton Permeability of

Thylakoid Membranes," J. Bioenerget. Biomembr. 14: 347-359 (1982)), peptidyl halomethyl ketones (chloromethyl or fluoromethyl ketones), peptidyl sulfonium salts, peptidyl acyloxymethyl ketones, derivatives and analogues of epoxides, such as E-64 (N-[N-(L-trans-carboxyoxiran-2-carbonyl)-L-leucylagmatine), E-64c (a derivative of E-64 in which the agmatine moiety is replaced by an isoamylamine moiety), E-64c ethyl ester, Ep-459 (an analogue of E-64 in which the agmatine moiety is replaced by a 1,4-diaminopropyl moiety), Ep-479 (an analogue of E-64 in which the agmatine moiety is replaced by a 1,7-diheptylamino moiety), Ep-460 (a derivative of Ep-459 in which the terminal amino group is substituted with a Z (benzyloxycarbonyl) group), Ep-174 (a derivative of E-64 in which the agmatine moiety is removed, so that the molecule has a free carboxyl residue from the leucine moiety), Ep-475 (an analogue of E-64 in which the agmatine moiety is replaced with a NH_2-$(CH_2)_2$-CH-$(CH_3)_2$ moiety), or Ep-420 (a derivative of E-64 in which the hydroxyl group is benzoylated, forming an ester, and the leucylagmatine moiety is replaced with isoleucyl-O-methyltyrosine), or peptidyl O-acyl hydroxamates (E Shaw, "Cysteinyl Proteases and Their Selective Inactivation), pp 271-347). Other inhibitors are known in the art.

Modification of other residues may also result in inhibition of the enzyme.

B. Screening Methods

Another aspect of the present invention is a method for screening a compound for anti-sortase-transamidase activity. This is an important aspect of the present invention, because it provides a method for screening for compounds that disrupt the sorting process and thus have potential antibiotic activity against Gram-positive bacteria.

In general, this method comprises the steps of: (1) providing an active fraction of sortase-transamidase enzyme; (2) performing an assay for sortase-transamidase activity in the presence and in the absence of the compound being screened; and (3) comparing the activity of the sortase-transamidase enzyme in the presence and in the absence of the compound.

The active fraction of sortase-transamidase enzyme can be a substantially purified sortase-transamidase enzyme preparation according to the present invention, but can be a less purified preparation, such as a partially purified particulate preparation as described below.

The enzymatic activity can be measured by the cleavage of a suitable substrate, such as the construct having the Staphylococcal Enterotoxin B (SEB) gene fused to the cell wall sorting signal of Staphylococcal Protein A (SPA). The cleavage can be determined by monitoring the molecular weight of the products by sodium dodecyl sulfate-polyacrylamide gel electrophoresis or by other methods.

One particularly preferred assay for sortase-transamidase activity is the following:

Staphylococcal soluble RNA (sRNA) is prepared from *S. aureus* by a modification of the technique of Zubay (G. Zubay, J. Mol. Biol. 4: 347-356 (1962)). An overnight culture of *S. aureus* is diluted 1:10 in TSB and incubated at 37°C for 3 hr. The cells are harvested by centrifugation at 6000 rpm for 15 min.

For every gram of wet cell pellets, 2 ml of 0.01 M magnesium acetate, 0.001 M Tris, pH 7.5 is used to suspend the pellets. The cell pellets are beaten by glass bead beater for 45 minutes in 5 minute intervals. The suspension is centrifuged twice at 2500 rpm for 5 minutes to remove the glass beads, then 0.5 ml phenol is added to the suspension. The suspension is vigorously shaken for 90 minutes at 4°C, and then centrifuged at 18,000 x g for 15 minutes. The nucleic acids in the top layer are precipitated by addition of 0.1 volume of 20% potassium acetate and 2 volumes of ethanol, then stored at 4°C for at least 36 hours. The precipitate is obtained by centrifugation at 5,000 x g for 5 minutes. Cold NaCl (1 ml) is added to the precipitate and stirred at 4°C for 1 hour. The suspension is centrifuged at 15,000 x g for 30 minutes. The sediments are washed with 0.5 ml of cold 1 M NaCl. The supernatants are combined and 2 volumes of ethanol is added to precipitate the tRNA. The precipitate is suspended in 0.1 ml of 0.2 M glycine, pH 10.3 and incubated for 3 hr at 37°C. This suspension is then made 0.4 M in NaCl and the RNA is precipitated by addition of 2 volumes of ethanol. The precipitate is dissolved in 0.7 ml of 0.3 M sodium acetate, pH 7.0. To this is slowly added 0.5 volume of isopropyl alcohol, with stirring. The precipitate is removed by centrifugation at 8,000 x g for 5 min. This precipitate is

redissolved in 0.35 ml of 0.3 M sodium acetate, pH 7.0. To this is added 0.5 volume of isopropyl alcohol, using the same procedure as above. The precipitate is also removed by centrifugation. The combined supernatants from the two centrifugations are treated further with 0.37 ml of isopropyl alcohol. The resulting precipitate is dissolved in 75 µl of water and dialyzed against water overnight at 4°C. This sRNA is used in the sortase-transamidase assay.

Particulate sortase-transamidase enzyme is prepared for use in the assay by a modification of the procedure of Chatterjee & Park (A.N. Chatterjee & J.T. Park, Proc. Natl. Acad. Sci. USA 51: 9-16 (1964)). An overnight culture of *S. aureus* OS2 is diluted 1:50 in TSB and incubated at 37°C for 3 hr. Cells are harvested by centrifugation at 6000 rpm for 15 minutes, and washed twice with ice-cold water. The cells are disrupted by shaking 7 ml of 1 3% suspension of cells in 0.05 M Tris-HCl buffer, pH 7.5, 0.1 mM $MgCl_2$, and 1 mM 2-mercaptoethanol with an equal volume of glass beads for 10-15 minutes in a beater. The glass beads are removed by centrifugation at 2000 rpm for 5 minutes. The crude extract is then centrifuged at 15,000 x g for 5 minutes. The supernatant is centrifuged again at 100,000 x g for 30 minutes. The light yellow translucent pellet is resuspended in 2 to 4 ml of 0.02 M Tris-HCl buffer, pH 7.5, containing 0.1 mM $MgCl_2$ and 1 mM 2-mercaptoethanol. This suspension represents the crude particulate enzyme and is used in the reaction mixture below.

The supernatant from centrifugation at 100,000 x g is passed through gel filtration using a Sephadex® G-25 agarose column (Pharmacia) to remove endogenous substrates. This supernatant is also used in the reaction mixture.

The complete reaction mixture contains in a final volume of 30 µl (M. Matsuhashi et al., Proc. Natl. Acad. Sci. USA 54: 587-594 (1965)): 3 µmol of Tris-HCl, pH 7.8; 0.1 µmol of $MgCl_2$; 1.3 µmol of KCl; 2.7 nmol of [^{3}H] glycine (200 µCi/µmol); 2 nmol of UDP-M-pentapeptide; 5 nmol of UDP-N-acetylglucosamine; 0.2 µmol of ATP; 0.05 µmol of potassium phosphoenolpyruvate; 2.05 µg of chloramphenicol; 5 µg of pyruvate kinase; 0.025 µmol of 2-mercaptoethanol; 50 µg of staphylococcal sRNA prepared as above; 4 µg (as protein) of supernatant as prepared above; 271 µg of particulate enzyme prepared as above; and 8 nmol of a synthesized soluble peptide (HHHHHHAQALEPTGEENPF) (SEQ ID NO: 32) as a substrate.

The mixture is incubated at 20°C for 60 minutes. The mixture is then heated at 100°C for 1 minute. The mixture is diluted to 1 ml and precipitated with 50 µl nickel resin, and washed with wash buffer (1% Triton X-100, 0.1% sodium dodecyl sulfate, 50 mM Tris, pH 7.5). The nickel resin beads are counted in a scintillation counter to determine ^{3}H bound to the beads.

The effectiveness of the compound being screened to inhibit the activity of the sortase-transamidase enzyme can be determined by adding it to the assay mixture in a predetermined concentration and determining the resulting degree of inhibition of enzyme activity that results. Typically, a dose-response curve is generated using a range of concentrations of the compound being screened.

The particulate enzyme preparation of sortase-transamidase employed in this protocol can be replaced with any other sortase-transamidase preparation, purified or crude, staphylococcal, recombinant, or from any other source from any other Gram-positive bacterium as described above.

The soluble peptide is captured in this embodiment by its affinity for nickel resin as a result of the six histidine residues. More than six histidine residues can be used in the peptide. As an alternative, the soluble peptide can be captured by an affinity resulting from other interactions, such as streptavidin-biotin, glutathione S-transferase-glutathione, maltose binding protein-amylose, and the like, by replacing the six histidine residues with the amino acid sequence that constitutes the binding site in the peptide and employing the appropriate solid phase affinity resin containing the binding partner. Suitable peptides can be prepared by solid phase peptide synthesis using techniques well known in the art, such as those described in M. Bodanszky, "Peptide Chemistry: A Practical Textbook" (2d ed., Springer-Verlag, Berlin, 1993). For example, if the glutathione S-transferase-glutathione interaction is used, the active site of glutathione S-transferase (D.B. Smith & K.S. Johnson, "Single-Step Purification of Polypeptides Expressed in *Escherichia coli* as Fusions with Glutathione S-Transferase," Gene 67: 31-40 (1988)) can be substituted for the six histidine residues, and glutathione can be bound to the solid support.

Alternatively, the soluble peptide can be released from the sortase by hydroxylaminolysis and then quantitated or monitored. The strong nucleophile hydroxylamine attacks thioester to form hydroxamate with carboxyl, thereby regenerating the enzyme sulfhydryl. Hydroxylaminolysis can be carried out in 50 mM Tris-HCl, pH 7.0 with a concentration of 0.1 M hydroxylamine for 60 minutes. The released peptide, for example, can be quantitated by mass spectroscopy or other methods.

IV. USE OF SORTASE-TRANSAMIDASE FOR PROTEIN AND PEPTIDE DISPLAY

A. Methods for Protein and Peptide Display

The sortase-transamidase enzyme of the present invention can also be used in a method of displaying a polypeptide on the surface of a gram-positive bacterium.

In general, a first embodiment of this method comprises the steps of: (1) expressing a polypeptide having a sorting signal at its carboxyl-terminal end as described above; (2) forming a reaction mixture including: (i) the expressed polypeptide; (ii) a substantially purified sortase-transamidase enzyme; and (iii) a Gram-positive bacterium having a peptidoglycan to which the sortase-transamidase can link the polypeptide; and (3) allowing the sortase-transamidase to catalyze a reaction that cleaves the polypeptide within the LPX_3X_4G motif of the sorting signal and covalently cross-links the amino-terminal portion of the cleaved polypeptide to the peptidoglycan to display the polypeptide on the surface of the Gram-positive bacterium.

In this method, the polypeptide having the sorting signal at its carboxy-terminal end need not be expressed in a Gram-positive bacterium; it can be expressed in another bacterial system such as *Escherichia coli* or *Salmonella typhimurium*, or in a eukaryotic expression system.

The other method for protein targeting and display relies on direct expression of the chimeric protein in a Gram-positive bacterium and the action of the sortase-transamidase on the expressed protein. In general, such a method comprises the steps of: (1) cloning a nucleic acid segment encoding a chimeric protein into a Gram-positive bacterium to generate a cloned chimeric protein including therein a carboxyl-terminal sorting signal as described above, the chimeric protein including the polypeptide to be displayed; (2) growing the bacterium into which the nucleic acid segment has been cloned to express the cloned chimeric protein to generate a chimeric protein including therein a carboxyl-terminal sorting signal; and (3) covalent binding of the chimeric protein to the cell wall by the enzymatic action of the sortase-transamidase involving cleavage of the chimeric protein within the LPX_3X_4G motif so that the protein is

displayed on the surface of the gram-positive bacterium in such a way that the protein is accessible to a ligand.

Typically, the Gram-positive bacterium is a species of *Staphylococcus*. A particularly preferred species of *Staphylococcus* is *Staphylococcus aureus*.

However, other Gram-positive bacteria such as *Streptococcus pyogenes*, other *Streptococcus* species, and Gram-positive bacteria of other genera can also be used.

Cloning the nucleic acid segment encoding the chimeric protein into the Gram-positive bacterium is performed by standard methods. In general, such cloning involves: (1) isolation of a nucleic acid segment encoding the protein to be sorted and covalently linked to the cell wall; (2) joining the nucleic acid segment to the sorting signal; (3) cloning by insertion into a vector compatible with the Gram-positive bacterium in which expression is to take place; and (4) incorporation of the vector including the new chimeric nucleic acid segment into the bacterium.

Typically, the nucleic acid segment encoding the protein to be sorted is DNA; however, the use of RNA in certain cloning steps is within the scope of the present invention.

When dealing with genes from eukaryotic organisms, it is preferred to use cDNA, because the natural gene typically contains intervening sequences or introns that are not translated. Alternatively, if the amino acid sequence is known, a synthetic gene encoding the protein to be sorted can be constructed by standard solid-phase oligodeoxyribonucleotide synthesis methods, such as the phosphotriester or phosphite triester methods. The sequence of the synthetic gene is determined by the genetic code, by which each naturally occurring amino acid is specified by one or more codons. Additionally, if a portion of the protein sequence is known, but the gene or messenger RNA has not been isolated, the amino acid sequence can be used to construct a degenerate set of probes according to the known degeneracy of the genetic code. General aspects of cloning are described, for example, in J. Sambrook et al., "Molecular Cloning: A Laboratory Manual" (2d ed., Cold Spring Harbor Laboratory Press, Cold Spring Harbor, New York, 1989); in B. Perbal, "A Practical Guide to Molecular Cloning" (2d ed., John Wiley & Sons, New York 1988), in S.L. Berger & A.R. Kimmel, "Guide to Molecular Cloning Techniques" (Methods in Enzymology, vol. 152, Academic Press, Inc., San Diego, 1987), and in

D.V. Goeddel, ed., "Gene Expression Technology" (Methods in Enzymology, vol. 185, Academic Press, Inc., San Diego, 1991).

Once isolated, DNA encoding the protein to be sorted is then joined to the sorting signal. This is typically accomplished through ligation, such as using *Escherichia coli* or bacteriophage T4 ligase. Conditions for the use of these enzymes are well known and are described, for example, in the above general references.

The ligation is done in such a way so that the protein to be sorted and the sorting signal are joined in a single contiguous reading frame so that a single protein is produced. This may, in some cases, involve addition or deletion of bases of the cloned DNA segment to maintain a single reading frame. This can be done by using standard techniques.

Cloning is typically performed by inserting the cloned DNA into a vector containing control elements to allow expression of the cloned DNA. The vector is then incorporated into the bacterium in which expression is to occur, using standard techniques of transformation or other techniques for introducing nucleic acids into bacteria.

One suitable cloning system for *S. aureus* places the cloned gene under the control of the BlaZRI regulon (P.Z. Wang et al., Nucl. Acids Res. 19:4000 (1991)). Vectors and other cloning techniques for use in *Staphylococcus aureus* are described in B. Nilsson & L. Abrahmsen, "Fusion to Staphylococcal Protein A," in Gene Expression Technology, supra, p.144-161.

If the chimeric protein is cloned under control of the BlaZRI regulon, expression can be induced by the addition of the β-lactam antibiotic methicillin.

Another aspect of the present invention is a polypeptide displayed on the surface of a Gram-positive bacterium by covalent linkage of an amino-acid sequence of LPX_3X_4 derived from cleavage of an LPX_3X_4G motif, as described above.

Yet another aspect of the present invention is a covalent complex comprising: (1) the displayed polypeptide; and (2) an antigen or hapten covalently cross-linked to the polypeptide.

B. Screening Methods

These polypeptides associated with the cell surfaces of Gram-positive bacteria can be used in various ways for screening. For example, samples of expressed proteins from an expression library containing expressed proteins on the surfaces of the cells can be used to screen for clones that express a particular desired protein when a labeled antibody or other labeled specific binding partner for that protein is available.

These methods are based on the methods for protein targeting and display described above.

A first embodiment of such a method comprises: (1) expressing a cloned polypeptide as a chimeric protein having a sorting signal at its carboxy-terminal end as described above; (2) forming a reaction mixture including: (i) the expressed chimeric protein; (ii) a substantially purified sortase-transamidase enzyme; and (iii) a Gram-positive bacterium having a peptidoglycan to which the sortase-transamidase can link the polypeptide through the sorting signal; (3) binding of the chimeric protein covalently to the cell wall by the enzymatic action of a sortase-transamidase expressed by the Gram-positive bacterium involving cleavage of the chimeric protein within the LPX_3X_4G motif so that the polypeptide is displayed on the surface of the Gram-positive bacterium in such a way that the polypeptide is accessible to a ligand; and (4) reacting the displayed polypeptide with a labeled specific binding partner to screen the chimeric protein for reactivity with the labeled specific binding partner.

The nucleic acid segment encoding the chimeric protein is formed by methods well known in the art and can include a spacer.

In the last step, the cells are merely exposed to the labeled antibody or other labeled specific binding partner, unreacted antibodies removed as by a wash, and label associated with the cells detected by conventional techniques such as fluorescence, chemiluminescence, or autoradiography.

A second embodiment of this method employs expression in a Gram-positive bacterium that also produces a sortase-transamidase enzyme. This method comprises: (1) cloning a nucleic acid segment encoding a chimeric protein into a Gram-positive bacterium to

generate a cloned chimeric protein including therein a carboxyl-terminal sorting signal as described above, the chimeric protein including the polypeptide whose expression is to be screened; (2) growing the bacterium into which the nucleic acid segment has been cloned to express the cloned chimeric protein to generate a chimeric protein including therein a carboxyl-terminal sorting signal; (3) binding the polypeptide covalently to the cell wall by the enzymatic action of a sortase-transamidase expressed by the Gram-positive bacterium involving cleavage of the chimeric protein within the LPX_3X_4G motif so that the polypeptide is displayed on the surface of the Gram-positive bacterium in such a way that the polypeptide is accessible to a ligand; and (4) reacting the displayed polypeptide with a labeled specific binding partner to screen the chimeric protein for reactivity with the labeled specific binding partner.

V. USE OF SORTED MOLECULES FOR DIAGNOSIS AND TREATMENT OF BACTERIAL INFECTIONS

Sorted molecules can also be used for the diagnosis and treatment of bacterial infections caused by Gram-positive bacteria. Antibiotic molecules or fluorescent or any other diagnostic molecules can be chemically linked to a sorted peptide segment, which may include a spacer as described above, and then can be injected into animals or humans. These molecules are then sorted by the sortase-transamidase so that they are covalently linked to the cell wall of the bacteria.

In general, these methods comprise: (1) conjugating an antibiotic or a detection reagent to a protein including therein a carboxyl-terminal sorting signal to produce a conjugate; and (2) introducing the conjugate to an organism infected with a Gram-positive bacterium in order to cause the conjugate to be sorted and covalently cross-linked to the cell walls of the bacterium in order to treat or diagnose the infection.

The antibiotic used can be, but is not limited to, a penicillin, ampicillin, vancomycin, gentamicin, streptomycin, a cephalosporin, amikacin, kanamycin, neomycin, paromomycin, tobramycin, ciprofloxacin, clindamycin, rifampin, chloramphenicol, or norfloxacin, or a derivative of these antibiotics.

The detection reagent is typically an antibody or other specific binding partner labeled with a detectable label, such as a radiolabel. Such methods are well known in the art and need not be described further here.

Accordingly, another aspect of the present invention is a conjugate comprising an antibiotic or a detection reagent covalently conjugated to a protein including therein a carboxyl-terminal sorting signal as described above to produce a conjugate.

Yet another aspect of the present invention is a composition comprising the conjugate and a pharmaceutically acceptable carrier.

In this context, the conjugates can be administered using conventional modes of administration, including, but not limited to, intravenous, intraperitoneal, oral, or intralymphatic. Other routes of administration can alternatively be used. Oral or intraperitoneal administration is generally preferred. The composition can be administered in a variety of dosage forms, which include, but are not limited to, liquid solutions or suspensions, tablets, pills, powders, suppositories, polymeric microcapsules or microvesicles, liposomes, and injectable or infusible solutions. The preferred form depends on the mode of administration and the quantity administered.

The compositions for administration preferably also include conventional pharmaceutically acceptable carriers and adjuvants known in the art such as human serum albumin, ion exchangers, alumina, lecithin, buffered substances such as phosphate, glycine, sorbic acid, potassium sorbate, and salts or electrolytes such as protamine sulfate. The most effective mode of administration and dosage regimen for the conjugates as used in the methods in the present invention depend on the severity and course of the disease, the patient's health, the response to treatment, the particular strain of bacteria infecting the patient, other drugs being administered and the development of resistance to them, the accessibility of the site of infection to blood flow, pharmacokinetic considerations such as the condition of the patient's liver and/or kidneys that can affect the metabolism and/or excretion of the administered conjugates, and the judgment of the treating physician. According, the dosages should be titrated to the individual patient.

VI. USE OF SORTED POLYPEPTIDES FOR PRODUCTION OF VACCINES

Additionally, the sorted polypeptides covalently crosslinked to the cell walls of Gram-positive bacteria according to the present invention have a number of uses. One use is use in the production of vaccines that can be used to generate immunity against infectious diseases affecting mammals, including both human and non-human mammals, such as cattle, sheep, and goats, as well as other animals such as poultry and fish. This invention is of special importance to mammals. The usefulness of these complexes for vaccine production lies in the fact that the proteins are on the surface of the cell wall and are accessible to the medium surrounding the bacterial cells, so that the antigenic part of the chimeric protein is accessible to the antigen processing system. It is well known that presenting antigens in particulate form greatly enhances the immune response. In effect, bacteria containing antigenic peptides on the surfaces linked to the bacteria by these covalent interactions function as natural adjuvants. Here follows a representative list of typical microorganisms that express polypeptide antigens against which useful antibodies can be prepared by the methods of the present invention:

(1) Fungi: *Candida albicans*, *Aspergillus fumigatus*, *Histoplasma capsulatum* (all cause disseminating disease), *Microsporum canis* (animal ringworm).

(2) Parasitic protozoa: (1) *Plasmodium falciparum* (malaria), *Trypanosoma cruzei* (sleeping sickness).

(3) Spirochetes: (1) *Borrelia bergdorferi* (Lyme disease), *Treponema pallidum* (syphilis), *Borrelia recurrentis* (relapsing fever), *Leptospira icterohaemorrhagiae* (leptospirosis).

(4) Bacteria: *Neisseria gonorrhoeae* (gonorrhea), *Staphylococcus aureus* (endocarditis), *Streptococcus pyogenes* (rheumatic fever), *Salmonella typhosa* (salmonellosis), *Hemophilus influenzae* (influenza), *Bordetella pertussis* (whooping cough), *Actinomyces israelii* (actinomycosis), *Streptococcus mutans* (dental caries), *Streptococcus equi* (strangles in horses), *Streptococcus agalactiae* (bovine mastitis), *Streptococcus anginosus* (canine genital infections).

(5) Viruses: Human immunodeficiency virus (HIV), poliovirus, influenza virus, rabies virus, herpes virus, foot and mouth disease virus, psittacosis virus, paramyxovirus, myxovirus, coronavirus.

Typically, the resulting immunological response occurs by both humoral and cell-mediated pathways. One possible immunological response is the production of antibodies, thereby providing protection against infection by the pathogen.

This method is not limited to protein antigens. As discussed below, non-protein antigens or haptens can be covalently linked to the C-terminal cell-wall targeting segment, which can be produced as an independently expressed polypeptide, either alone, or with a spacer at its amino-terminal end. If a spacer at the amino-terminal end is used, typically the spacer will have a conformation allowing the efficient interaction of the non-protein antigen or hapten with the immune system, most typically a random coil or α-helical form. The spacer can be of any suitable length; typically, it is in the range of about 5 to about 30 amino acids; most typically, about 10 to about 20 amino acids. In this version of the embodiment, the independently expressed polypeptide, once expressed, can then be covalently linked to the hapten or non-protein antigen. Typical non-protein antigens or haptens include drugs, including both drugs of abuse and therapeutic drugs, alkaloids, steroids, carbohydrates, aromatic compounds, including many pollutants, and other compounds that can be covalently linked to protein and against which an immune response can be raised.

Alternatively, a protein antigen can be covalently linked to the independently expressed cell-wall targeting segment or a cell-wall targeting segment including a spacer.

Many methods for covalent linkage of both protein and non-protein compounds to proteins are well known in the art and are described, for example, in P. Tijssen, "Practice and Theory of Enzyme Immunoassays" (Elsevier, Amsterdam, 1985), pp. 221-295, and in S.S. Wong, "Chemistry of Protein Conjugation and Cross-Linking" (CRC Press, Inc., Boca Raton, FL, 1993).

Many reactive groups on both protein and non-protein compounds are available for conjugation.

For example, organic moieties containing carboxyl groups or that can be carboxylated can be conjugated to proteins via the mixed anhydride method, the carbodiimide method, using dicyclohexylcarbodiimide, and the N-hydroxysuccinimide ester method.

If the organic moiety contains amino groups or reducible nitro groups or can be substituted with such groups, conjugation can be achieved by one of several techniques. Aromatic amines can be converted to diazonium salts by the slow addition of nitrous acid and then reacted with proteins at a pH of about 9. If the organic moiety contains aliphatic amines, such groups can be conjugated to proteins by various methods, including carbodiimide, tolylene-2,4-diisocyanate, or malemide compounds, particularly the N-hydroxysuccinimide esters of malemide derivatives. An example of such a compound is 4-(N-maleimidomethyl)-cyclohexane-1-carboxylic acid. Another example is m-maleimidobenzoyl-N-hydroxysuccinimide ester. Still another reagent that can be used is N-succinimidyl-3-(2-pyridyldithio) propionate. Also, bifunctional esters, such as dimethylpimelimidate, dimethyladipimidate, or dimethylsuberimidate, can be used to couple amino-group-containing moieties to proteins.

Additionally, aliphatic amines can also be converted to aromatic amines by reaction with p-nitrobenzoylchloride and subsequent reduction to a p-aminobenzoylamide, which can then be coupled to proteins after diazotization.

Organic moieties containing hydroxyl groups can be cross-linked by a number of indirect procedures. For example, the conversion of an alcohol moiety to the half ester of succinic acid (hemisuccinate) introduces a carboxyl group available for conjugation. The bifunctional reagent sebacoyldichloride converts alcohol to acid chloride which, at pH 8.5, reacts readily with proteins. Hydroxyl-containing organic moieties can also be conjugated through the highly reactive chlorocarbonates, prepared with an equal molar amount of phosgene.

For organic moieties containing ketones or aldehydes, such carbonyl-containing groups can be derivatized into carboxyl groups through the formation of O-(carboxymethyl) oximes. Ketone groups can also be derivatized with p-hydrazinobenzoic acid to produce carboxyl groups that can be conjugated to the specific binding partner as described above.

Organic moieties containing aldehyde groups can be directly conjugated through the formation of Schiff bases which are then stabilized by a reduction with sodium borohydride.

One particularly useful cross-linking agent for hydroxyl-containing organic moieties is a photosensitive noncleavable heterobifunctional cross-linking reagent, sulfosuccinimidyl 6-[4′-azido-2′-nitrophenylamino] hexanoate. Other similar reagents are described in S.S. Wong, "Chemistry of Protein Conjugation and Cross-Linking," supra.

Other cross-linking reagents can be used that introduce spacers between the organic moiety and the specific binding partner.

These methods need not be described further here.

VII. PRODUCTION OF SUBSTANTIALLY PURIFIED SORTASE-TRANSAMIDASE ENZYME

Another aspect of the present invention is methods for the production of substantially purified sortase-transamidase enzyme.

A. Methods Involving Expression of Cloned Gene

One method for the production of substantially purified sortase-transamidase enzyme involves the expression of the cloned gene, preferably the *srt*A gene. The isolation of the nucleic acid segment or segments encoding the sortase-transamidase enzyme is described above; these nucleic acid segment or segments are then incorporated into a vector and then use to transform a host in which the enzyme can be expressed. In one alternative, the host is a Gram-positive bacterium.

The next step in this alternative is expression in a Gram-positive bacterium to generate the cloned sortase-transamidase enzyme. Expression is typically under the control of various control elements associated with the vector incorporating the DNA encoding the sortase-transamidase gene, such as the coding region of the *srt*A gene; such elements can include promoters and operators, which can be regulated by proteins such as repressors. The conditions required for expression of cloned proteins in gram-positive bacteria, particularly *S. aureus*, are

well known in the art and need not be further recited here. An example is the induction of expression of lysostaphin under control of the BlaZRI regulon induced by the addition of methicillin.

When expressed in *Staphylococcus aureus*, the chimeric protein is typically first exported with an amino-terminal leader peptide, such as the hydrophobic signal peptide at the amino-terminal region of the cloned lysostaphin of Recsei et al. (P. Recsei et al., "Cloning, Sequence, and Expression of the Lysostaphin Gene from *Staphylococcus simulans*," Proc. Natl. Acad. Sci. USA 84:1127-1131 (1987)).

Alternatively, the cloned nucleic acid segment encoding the sortase-transamidase enzyme can be inserted in a vector that contains sequences allowing expression of the sortase-transamidase in another organism, such as *E. coli* or *S. typhimurium*. A suitable host organism can then be transformed or transfected with the vector containing the cloned nucleic acid segment. Expression is then performed in that host organism.

The expressed enzyme is then purified using standard techniques. Techniques for the purification of cloned proteins are well known in the art and need not be detailed further here. One particularly suitable method of purification is affinity chromatography employing an immobilized antibody to sortase. Other protein purification methods include chromatography on ion-exchange resins, gel electrophoresis, isoelectric focusing, and gel filtration, among others.

One particularly useful form of affinity chromatography for purification of cloned proteins, such as sortase-transamidase, as well as other proteins, such as glutathione S-transferase and thioredoxin, that have been extended with carboxyl-terminal histidine residues, is chromatography on a nickel-sepharose column. This allows the purification of a sortase-transamidase enzyme extended at its carboxyl terminus with a sufficient number of histidine residues to allow specific binding of the protein molecule to the nickel-sepharose column through the histidine residues. The bound protein is then eluted with imidazole. Typically, six or more histidine residues are added; preferably, six histidine residues are added. One way of adding the histidine residues to a cloned protein, such the sortase-transamidase, is through PCR with a primer that includes nucleotides encoding the histidine residues. The histidine codons are

CAU and CAC expressed as RNA, which are CAT and CAC as DNA. Amplification of the cloned DNA with appropriate primers will add the histidine residues to yield a new nucleic acid segment, which can be recloned into an appropriate host for expression of the enzyme extended with the histidine residues.

B. Other Methods

Alternatively, the sortase-transamidase can be purified from Gram-positive bacteria by standard methods, including precipitation with reagents such as ammonium sulfate or protamine sulfate, ion-exchange chromatography, gel filtration chromatography, affinity chromatography, isoelectric focusing, and gel electrophoresis, as well as other methods known in the art.

Because the sortase-transamidase is a cysteine protease, one particularly useful method of purification involves covalent chromatography by thiol-disulfide interchange, using a two-protonic-state gel containing a 2-mercaptopyridine leaving group, such as Sepharose 2B-glutathione 2-pyridyl disulfide or Sepharose 6B-hydroxypropyl 2-pyridyl disulfide. Such covalent chromatographic techniques are described in K. Brocklehurst et al., "Cysteine Proteases," in New Comprehensive Biochemistry, Volume 16: Hydrolytic Enzymes (A. Neuberger & K. Brocklehurst, eds., Elsevier, New York, 1987), ch. 2, pp. 39-158.

VIII. FURTHER APPLICATIONS OF SORTASE-TRANSAMIDASE

A. Production of Antibodies

Antibodies can be prepared to the substantially purified sortase-transamidase of the present invention, whether the sortase-transamidase is purified from bacteria or produced from recombinant bacteria as a result of gene cloning procedures. Because the substantially purified enzyme according to the present invention is a protein, it is an effective antigen, and antibodies can be made by well-understood methods such as those disclosed in E. Harlow & D. Lane, "Antibodies: A Laboratory Manual" (Cold Spring Harbor Laboratory, 1988). In general, antibody preparation involves immunizing an antibody-producing animal with the protein, with

or without an adjuvant such as Freund's complete or incomplete adjuvant, and purification of the antibody produced. The resulting polyclonal antibody can be purified by techniques such as affinity chromatography.

Once the polyclonal antibodies are prepared, monoclonal antibodies can be prepared by standard procedures, such as those described in Chapter 6 of Harlow & Lane, supra.

B. Derivatives for Affinity Chromatography

Another aspect of the present invention is derivatives of the cloned, substantially purified sortase-transamidase of the present invention extended at its carboxyl terminus with a sufficient number of histidine residues to allow specific binding of the protein molecule to a nickel-sepharose column through the histidine residues. Typically, six or more histidine residues are added; preferably, six histidine residues are added.

The histidine residues can be added to the carboxyl terminus through PCR cloning as described above.

This invention is further described by means of the following example. This Example is for illustrative purposes only, and are not to be construed as limiting the scope of the invention in any manner.

Example 1

Identification of a Staphylococcal Mutant Defective in Cell Wall Sorting

Generation of temperture sensitive (ts) mutants through chemical mutagenesis

Cell wall sorting mutants were created and isolated from a population of conditional lethal mutants of *S. aureus* strain OS2. Staphylococci were mutagenized with nitrosoguanidine and colonies were formed by plating at 30°C. Bacteria were streaked and incubated at 30°C and 42°C to identify mutants that are temperature sensitive for growth (ts). A collection of one thousand ts mutants was transformed with pSEB-SPA$_{490-524}$ (O. Schneewind, D. Mihaylova-Petkov, P. Model, *EMBO* **12**, 4803 (1993)), specifying a reporter protein for measurements of surface protein anchoring. The SEB-SPA$_{490-524}$ precursor (P1) is exported from the cytoplasm and its NH_2-terminal leader peptide removed to generate the P2 intermediate (Figure 2A). The P2 precursor is the substrate for sortase, which cleaves the polypeptide between the threonine and the glycine of the LPXTG motif and generates mature, anchored surface protein (M). When analyzed by labeling wild-type staphylococci with [^{35}S]methionine for 5 min, cleavage of P1 precursor is faster than that of the P2 species, yielding a ratio of P1 (5%), P2 (19%), and M(76%) concentration (Figure 2B). This assay was employed to screen one thousand ts mutants and two strains were identified that accumulated P2 precursor at 47% (SM317) and 26% (SM329), respectively (Figure 2B). To examine the sorting reaction further, mutant and wild-type staphylococci were subjected to pulse-chase analysis (Figure 2C). *S. aureus* OS2 (wild-type) cleaved and anchored the P1 precursor within 2 min. The sorting reaction in strain SM317 was severely reduced as cleavage and cell wall anchoring of pulse-labeled P2 required more than 10 min. Strain SM329 displayed only a weak defect and P2 processing required 3 min (Figure 2C). When examined by pulse-labeling staphylococci grown in minimal medium, SM329 displayed a much more severe defect in cell wall sorting.

Anchor structure of surface proteins in the mutant strain SM317

To examine whether the mutant strains SM317 and SM329 are defective in the synthesis of bacterial cell wall, two tests were performed. Lysostaphin is a bacteriolytic enzyme that cuts the pentaglycine crossbridges of the staphylococcal cell wall predominantly at the central glycine residue (C. A. Schindler and V. T. Schuhardt, *Proc. Natl. Acad. Sci. USA* **51**, 414 (1964); B. L. M. de Jonge, Y. S. Chang, D. Gage, A. Tomasz, *J. Biol. Chem.* **267**, 11248 (1992)). As reported previously, *fem* mutants display resistance to this bacteriocin and grow even in the presence of large amounts of lysostaphin (U. Kopp, M. Roos, J. Wecke, H. Labischinski, *Microb. Drug Resist.* **2**, 29 (1996)). Strains SM317 and SM329 were sensitive to lysostaphin at concentrations that also inhibited growth of wild-type staphylococci, indicating that the sorting defect in SM317 is not caused by a mutationally altered cell wall crossbridge. To measure bacterial cell wall synthesis, staphylococci were grown in minimal medium and labeled with [^{3}H]lysine and [^{3}H]leucine (D. Boothby, L. Daneo-Moore, G. D. Shockman, *Anal. Biochem.* **44**, 645 (1971)). As lysine, but not leucine, is a component of the bacterial cell wall, the ratio of [^{3}H]lysine/[^{3}H]leucine incorporation into acid precipitable and protease resistant murein polymer is a measure for cell wall synthesis (D. Boothby, L. Daneo-Moore, G. D. Shockman, *Anal. Biochem.* **44**, 645 (1971)). Wild-type staphylococci displayed a ratio of 30, while the addition of vancomycin to the culture medium reduced the ratio of incorporated lysine/leucine to 1.5 (20 fold inhibition). Strains SM317 and SM329 displayed a ratio of 18 and 19 (1.6 fold less than wild-type cells), suggesting that the accumulation of P2 precursor in the mutant SM317 is not caused by a defect in cell wall synthesis.

The cell wall anchor structure of surface protein in strain SM317 was determined (Figure 3). Plasmid pHTT4 specifying the reporter protein SEB-MH$_6$-CWS was transformed into *S. aureus* SM317 (H. Ton-That, K. F. Faull, O. Schneewind, *J. Biol. Chem.* **272**, 22285 (1997)). The staphylococcal cell wall was purified and digested with mutanolysin, a muramidase that hydrolyzes the glycan strands (K. Yokogawa, *et al.*, *Antimicrob. Agents Chemother.* **6**, 156 (1974)). Mutanolysin-released surface protein was purified by chromatography on Ni-NTA and cleaved at methionine residues with cyanogen bromide (H. Ton-That, K. F. Faull, O.

Schneewind, *J. Biol. Chem.* **272**, 22285 (1997)). COOH-terminal peptides bearing cell wall anchor structures were purified by a second affinity chromatography step and analyzed by MALDI-MS (Figure 3B). A series of ion signals with regularly spaced mass increments was revealed, measurements that are consistent with one, two, three, four, five and six peptidoglycan subunits linked to the COOH-terminal threonine of surface protein. Ion signals of muanolysin-solubilized anchor peptides were explained as H_6AQALPET-Gly$_5$ linked to cell wall tetrapeptide (predicted mass 2235; observed 2236), pentapeptide (predicted mass 2306; observed 2306), N,O6-diacetylMurNac-GlcNac tetrapeptide (predicted mass 2755, observed 2756), N,O6-diacetylMurNac-GlcNac pentapeptide (predicted mass 2826, observed 2826), murein-tetrapeptide-murein-pentapeptide (predicted mass 3991, observed 3995), (murein-tetrapeptide)$_2$-murein-pentapeptide (predicted mass 5194; observed 5196), (murein-tetrapeptide)$_4$ (predicted mass 6285 observed 6285), (murein-tetrapeptide)$_4$-murein-pentapeptide (predicted mass 7581; observed 7583), (murein-tetrapeptide)$_5$-murein-pentapeptide (predicted mass 8783; observed 8784). If surface protein is tethered to cross-linked peptidoglycan of strain SM317, digestion of muramidase-solubilized anchor peptides with *f*11 hydrolase should produce anchor peptide linked to murein tetrapeptide and disaccharide-tetrapeptide (H. Ton-That, K. F. Faull, O. Schneewind, *J. Biol. Chem.* **272**, 22285 (1997); W. W. Navarre, H. Ton-That, K. F. Faull, O. Schneewind, *J. Biol. Chem.* **274**, in press (1999)) (Figure 3). This was tested and the doubly digested anchor peptides generated ion signals at *m/z* 2236 [L-Ala-D-iGln-L-Lys(NH_2-H_6AQALPET-Gly$_5$)-D-Ala, predicted mass 2235], 2714 [MurNac(L-Ala-D-iGln-L-Lys(NH_2-H_6AQALPET-Gly$_5$)-D-Ala)-GlcNac, predicted mass 2713] and 2756 [O6-acetyl-MurNac(L-Ala-D-iGln-L-Lys(NH_2-H_6AQALPET-Gly$_5$)-D-Ala)-GlcNac, predicted mass 2756] (Figure 3C). Thus, surface proteins of *S. aureus* SM317 are tethered to cross-linked peptidoglycan in a manner that is indistinguishable from the anchor structure of polypeptides in wild-type staphylococci (W. W. Navarre, H. Ton-That, K. F. Faull, O. Schneewind, *J. Biol. Chem.* **273**, 29135 (1998)). These results suggest that the accumulation of P2 precursor in strain SM317 is likely caused by a defect in sortase.

Screening for the Sortase Gene

Over-expression of sortase from a multi-copy plasmid should reduce the concentration of P2 in both wild-type and mutant staphylococci. A plasmid library of two thousand 3-5 kb random *S. aureus* OS2 chromosomal DNA insertions was screened for sequences that caused a reduction in the concentration of P2 precursor in strain SM317. Two plasmids, pGL1631 and pGL1834, answered this screen (Figure 4). Transformation with pGL1834 reduced the P2 concentration in strain SM317 from 44% to 9%, in strain SM329 from 26% to 12%, and in wild-type *S. aureus* OS2 from 17% to 8%. When measured by pulse-chase analysis, *S. aureus* OS2 (pGL1834) displayed a rapidly increased processing of P2 precursors, a phenotype that was also observed in strains SM317 and SM329 (Figure 4C). DNA sequencing revealed that pGL1631 and pGL1834 contained staphylococcal chromosomal DNA insertions with identical overlapping sequences. The DNA sequence sufficient to promote a reduction in P2 concentration was mapped to a gene which was named *srtA* (surface protein sorting A) (Figure 5).

The *srtA* gene

The *srtA* gene (SEQ. ID NO. 2) specifies a polypeptide chain of 206 amino acids (Figure 6; SEQ. ID. NO. 3). A sequence of 18 hydrophobic amino acids near the NH_2-terminus suggests the presence of a signal peptide/membrane anchor sequence. This feature is consistent with the notion that cell wall anchoring occurs on the cell surface, after polypeptide substrates bearing an LPXTG motif have been translocated across the cytoplasmic membrane. Another property of the *srtA* gene consistent with its function as sortase is the presence of codon 184 specifying cysteine. As the cell wall sorting reaction is sensitive to methanethiosulfonate, a reagent that forms disulfide with sulfhydryl (D.J. Smith, E.T. Maggio, G.L. Kenyon, *Biochemistry* **14**, 764 (1975)), the presence of a cysteine must be a conserved feature of sortase homologues.

Many, if not all, Gram-positive pathogens display proteins on their surface via a sorting signal mediated mechanism (W. W. Navarre and O. Schneewind, *Microbiol. Mol. Biol.*

Rev. **63**, 174 (1999)). Thus, if the *srtA* gene specifies sortase, homologous genes should be found in the genomes of other Gram-positive pathogens. Chromosomal DNA sequences of *Enterococcus faecalis*, *Staphylococcus aureus*, *Streptococcus pyogenes*, *Streptococcus pneumoniae*, and *Streptococcus mutans* were searched and the presence of *srtA* genes revealed (Figure 7). Database searches also identified sequences homologous to *srtA* in *Bacillus subtilis* and *Actinomyces naeslundii*. All *srtA* homologues displayed absolute conservation of the cysteine and striking conservation of the peptide sequences surrounding it (Figure 7). *S. pneumoniae* harbors more than one *srtA* homologue which we have named *srtB* and *srtC*, respectively. The *srtA* like genes of *E. faecalis* and *A. naeslundii* are immediately adjacent to structural genes specifying surface proteins with a COOH-terminal sorting signal. The presence of a *srtA* homologue in the chromosome of *B. subtilis* is surprising as LPXTG motif containing sorting signals have not yet been identified in this organism. One of the *srtA* homologues in *A. naeslundii*, previously designated *orf365*, has been mutated, which abolished fimbrial assembly of mutant Actinomyces (M. K. Yeung, J. A. Donkersloot, J. O. Cisar, P. A. Ragsdale, *J. Bacteriol.* **66**, 1482 (1998)). Actinomyces fimbriae are composed of protein subunits bearing LPXTG motifs (M. K. Yeung and J. O. Cisar, *J. Bacteriol.* **172**, 2462 (1990)), however the mechanism of fimbrial assembly (polymerization) is not yet understood.

The *srtA* gene in strain SM317

To examine whether the defect in cell wall sorting of *S. aureus* SM317 is caused by a mutation in the *srtA* gene, corresponding sequences were PCR amplified from the chromosomal DNA of *S. aureus* OS2 and SM317. When cloned into a multi-copy vector and transformed into *S. aureus* SM317, the *srtA* gene amplified from wild-type staphylococci reduced the P2 concentration from 44% to 12%, while the same gene amplified from the chromosomal DNA of *S.aureus* SM317 did not reduce the P2 concentration of the parent strain (Figure 4B). Thus, the *srtA* gene is defective in strain SM317 and DNA sequencing identified mutations in codons 35 and 180. The expression of wild-type *srtA* in SM317 in the ts phenotype of the mutant strain was examined. Multi-copy expression of *srtA* (pGL1894) allowed growth of

SM317 at 42°C albeit at a rate that was less than that observed for wild-type staphylococci. This result suggests that the conditional lethal phenotype of *S. aureus* SM317 is not only caused a mutation in the *srtA* gene. Expression of plasmid encoded wild-type *srtA* did not alter the ts growth phenotype of *S. aureus* SM329.

Sortase and the cell wall sorting reaction

The *srtA* gene was isolated as a multi-copy suppressor of P2 precursor accumulation, a scheme that should only be answered by the gene for sortase. Only one gene (*srtA*) from a library of two thousand plasmid transformants bearing random 3-5 kb chromosomal DNA insertions was observed this screen. Additional observations show SrtA protein catalyzes the *in vitro* transpeptidation of substrates bearing an LPXTG motif, thereby demonstrating that SrtA displays sortase activity. Purified SrtA protein can be used for the screening of compounds that inhibit sortase. Such compounds may be useful for the treatment of human infections caused by Gram-positive bacteria.

Materials and Methods

Mutagenesis of S. aureus Strain OS2

Staphylococci (1 x 10^{12} cfu) were treated with 0.2 mg/ml N-methyl-N'-nitro-N-nitrosoguanidine for 45 min at 30°C and mutagenesis was quenched by the addition of 2 volumes of 100 mM sodium phosphate, pH 7.0. Approximately 80% of the mutagenized population was killed and the mutational frequency of rifampicin resistant *rpoB* mutations was increased to 1.2 x 10^{-4}. Temperature sensitive mutants were selected by growing the mutagenized population in tryptic soy broth at 42°C and treating with 8 μg/ml penicillin G for two hours, a selection that was repeated twice. Colonies were formed at 30°C, streaked on tryptic soy agar and examined for growth at 42°C.

Transformation of Competent Cells

Staphylococci were grown in tryptic soy broth supplemented with chloramphenicol (10 mg/ml) or tetracycline (2 mg/ml) at 30°C until OD_{660} 0.6. Cells were incubated at 42°C for 20 min, sedimented by centrifugation at 15,000 x *g* for 3 minutes and washed with 1 ml of prewarmed minimal medium [Schneewind, O., Model, P., Fischetti, V.A. (1992) Cell 70, 267]. Staphylococci were labeled with 50 mCi of [^{35}S]-Promix (Amersham) for 5 minutes and surface protein processing quenched by the addition of 75 ml 100% TCA. The TCA precipitates were collected by centrifugation, washed in acetone and dried under vacuum. Samples were suspended in 1 ml of 0.5 M Tris-HCl, pH 7.0 and staphylococcal peptidoglycan was digested by adding 50 ml 2 mg/ml lysostaphin (AMBI Pharmaceuticals) for 1 hour at 37°C. Proteins were again precipitated with TCA, washed with acetone and, after immunoprecipitation with a-SEB, were analyzed by 14% SDS-PAGE and PhosphorImager.

Pulse-Chase Screen of Mutants

Staphylococci were grown as described above and 5 ml were labeled with 500 mCi of [^{35}S]-Promix (Amersham) for 45 seconds. Incorporation of radioactivity was quenched by adding 50 ml chase (100 mg/ml casamino acids, 20 mg/ml methionine and cysteine). At timed intervals after the addition of the chase, 1 ml aliquots were removed and protein was precipitated by the addition of 75 ml 100% TCA. Sample preparation followed the same steps as described above.

DNA Sequencing

The DNA insertions pf pGL1631 and 1834 were mapped and sequenced by synthesizing oligonucleotide primers that annealed to sequenced template DNA 500 nucleotides apart. The primers for the amplification of *srtA* from the chromosomal DNA of *S. aureus* strains OS2 and SM317 were 5'-AAGGATTCAAAAGGAGCGGTATACATTGC-3' (SEQ ID NO. 32) and 5'-AAGGATCCTACCTTTTCCTCTAGCTGAAC-3' (SEQ ID NO. 33).

EXAMPLE 2

Inhibitors of Cell Wall Sorting

To study the effects of antibiotc cell wall synthesis inhibitors interfered with the anchoring of surface proteins, the activity of several inhibitors were examined in a Gram-positive bacteria sorting assay. A search for chemical inhibitors of the sorting reaction identified methanethiosulfonates and p-hydroxymercuribenzoic acid. Thus, sortase, the enzyme proposed to cleave surface proteins at the LPXTG motif, appears to be a sulfhydryl containing enzyme that utilizes peptidoglycan precursors but not assembled cell wall as a substrate for the anchoring of surface protein.

In order to identify compounds that interfere with the anchoring of surface proteins a reporter protein Seb-Spa490-524 which, when expressed in *S. aureus* OS2 cells, is synthesized as a precursor in the cytoplasm and initiated into the secretory pathway by an NH_2-terminal leader peptide (P1 precursor) was utilized (Schneewind, O., Mihaylova-Petkov, D. and Model, P. (1993) *EMBO* **12**, 4803-4811). After signal peptide cleavage, the P2 precursor bearing a COOH-terminal sorting signal serves as a substrate for sortase, an enzyme that cleaves between the threonine and the glycine of the LPXTG motif (Navarre, W. W. and Schneewind, O. (1994) *Mol. Microbiol.* **14**, 115-121). Amide linkage of the carboxyl of threonine to the cell wall crossbridge generates mature, anchored surface protein (M) (Schneewind, O., Fowler, A. and Faull, K. F. (1995) *Science* **268**, 103-106). Surface protein processing was investigated by pulse-labeling polypeptides with [^{35}S]methionine. During the pulse, all three species, P1 and P2 precursors as well as mature Seb-Spa490-524 can be detected (Figure 8B). Within 1 min after the addition of the chase, most pulse-labeled surface protein was converted to the mature, anchored species. Surface protein anchoring was complete 3 min after the quenching of [^{35}S]methionine incorporation.

Sodium azide is an inhibitor of SecA, an essential component of the secretory pathway in bacteria (Oliver, D. B., Cabelli, R. J., Dolan, K. M. and Jarosik, G. P. (1990) *Proc. Natl. Acad. Sci. USA* **87**, 8227-8231). Addition of 5 mM sodium azide to staphylococcal cultures

5 min prior to pulse-labeling significantly reduced protein export and led to the accumulation of leader peptide bearing P1 precursor (Schneewind, O., Model, P. and Fischetti, V. A. (1992) *Cell* **70**, 267-281). Methanethiosulfonates react with sulfhydryl (Akabas, M. H. and Karlin, A. (1995) *Biochemistry* **34**, 12496-12500) and one of these compounds, [2-(trimethylammonium) ethyl]methanethiosulfonate) (MTSET) prevented incorporation of [^{35}S]methionine by staphylococci. However, when added 15 seconds after the beginning of the pulse, MTSET interfered with the cleavage of sorting signals at the LPXTG motif, while the Sec-dependent export of P1 precursor remained unaltered. This result revealed that sortase must harbor a sulfhydryl that is necessary for enzymatic cleavage at LPXTG bearing sorting signals.

Sortase's requirement on sulfhydryl for enzymatic activity was tested, by the addition of other sulfhydryl reagents and analysis of inhibition of the cleavage of sorting signals at the LPXTG motif. MTSES, another methanethiosulfonate also interfered with sorting albeit not as effectively as MTSET (Table I). pHMB, an organic mercurial known to inhibit cysteine proteases, also displayed an inhibitory effect, whereas alkylating reagents such as N-ethylmaleimide, iodoacetate and iodoacetamide did not (Creighton, T. E. (1993) *Proteins*. W.H. Freeman and Company, New York.). Sulfhydryl reducing agents, i.e. dithiothreitol and mercaptoethanol, did not affect the sorting reaction. Neither PMSF, which reacts with hydroxyl (Creighton, T. E. (1993) *Proteins*. W.H. Freeman and Company, New York), nor treatment with the divalent cation chelator EDTA interfered with cell wall sorting, indicating that sortase likely does not require divalent cations or hydroxyl for cleavage and anchoring of surface protein.

Antibiotic inhibition of bacterial cell wall synthesis and cell wall sorting

To examine the effect of known antibiotics on cell wall sorting three compounds, penicillin, vancomycin and moenomycin were used. *S. aureus* OS2 (pSeb-Spa$_{490-524}$) was grown in minimal medium until A_{600} of 0.3, treated with 10 μg/ml of either penicillin, vancomycin, or moenomycin and incubated for an additional 5 hours (Figure 9A). At 30 min intervals during this experiment, aliquots were withdrawn for measurements of surface protein sorting and cell wall synthesis. The effect of antibiotics on the rate of bacterial cell wall

synthesis was determined as the ratio of [^{3}H]lysine/[^{3}H]leucine label incorporated into acid precipitable, pronase resistant peptidoglycan. Lysine is a component of peptidoglycan, whereas leucine is not. Hence, the ratio of incorporation of these two amino acids is a measure for cell wall synthesis. Surface protein anchoring was measured by pulse-labeling and quantified as the ratio between the concentration of P2 precursor [P2] and mature, anchored Seb-Spa$_{490-524}$ [M].

Addition of vancomycin, penicillin or moenomycin reduced the growth rate of staphylococci as compared to a mock treated control. While the rate of cell wall sorting precursor cleavage remained constant during the growth of mock treated staphylococci, the addition of vancomycin led to a steady accumulation of P2 precursor, indicating that this compound caused a reduction of the sorting reaction. A similar, albeit weaker effect was observed when moenomycin was added to staphylococcal cultures. In contrast, penicillin G did not alter the rate of cell wall sorting. As expected, all three antibiotics diminished the rate of peptidoglycan synthesis (Table II). Together these data revaled that vancomycin and moenomycin cause a reduction in the rate of cell wall sorting, while penicillin had no effect on surface protein anchoring.

Cell wall sorting in staphylococcal protoplasts

Previous work revealed that protoplasts, generated by muralytic digestion of staphylococci or penicillin selection of streptococcal L forms, secreted surface protein into the surrounding medium (van de Rijn, I. and Fischetti, V. A. (1981) *Infect. Immun.* **32**, 86-91; Movitz, J. (1976) *Eur. J. Biochem.* **68**, 291-299). This can be explained in two ways. Either the C-terminal sorting signals cannot retain surface proteins in the envelope of protoplasts or the presence of intact, assembled cell wall is not required to cleave sorting signals at their LPXTG motif. To distinguish between these possibilities, the surface protein anchoring in intact bacteria and staphylococcal protoplasts was measured (Figure 10). Wild-type staphylococci cleaved the Seb-Cws-BlaZ precursor to generate the mature, anchored NH_2-terminal Seb and COOH-terminal, cytoplasmic BlaZ fragments (Navarre, W. W. and Schneewind, O. (1994) *Mol. Microbiol.* **14**, 115-121). When tested in staphylococcal protoplasts generated by lysostaphin-

digestion of the cell wall, precursor cleavage occurred similar to whole cells, indicating that the presence of mature, assembled cell wall is not required for cleavage of sorting signals. Unique sorting products in protoplasts that migrated more slowly than mature, anchored Seb (see arrow in Figure 10B) were observed. As these species were immunoprecipitated with a-Seb but not with a-BlaZ, they likely represent products of the sorting reaction. The COOH-terminal anchor structure of these protoplast species are distinct from those generated by lysostaphin-digestion (three glycyl attached to the carboxyl of threonine), as they migrated more slowly on SDS-PAGE than lysostaphin-released Seb.

To examine whether all cleaved Seb fragments were released into the extra-cellular medium, pulse-labeled protoplasts were sedimented by centrifugation and separated from the extra-cellular medium in the supernatant. All Seb-Cws-BlaZ precursor and COOH-terminal BlaZ cleavage fragment sedimented with the protoplasts. In contrast, NH_2-terminal Seb fragments that migrated at the same speed as Seb released by lysostaphin-digestion from the cell wall of intact staphylococci were soluble in the culture medium. Some, but not all, of the more slowly migrating Seb species sedimented into the pellet, suggesting that these products of the sorting reaction may be attached to protoplast membranes. No precursor cleavage was observed for Seb-Cws$_{DLPXTG}$-BlaZ in either whole cells or staphylococcal protoplasts.

Materials and Methods

Bacterial Strains and Plasmids

Plasmids pSeb-Spa490-524(3), pSeb-Csw-BlaZ, and pSeb-Cws$_{DLPXTG}$-BlaZ (Navarre, W. W. and Schneewind, O. (1994) *Mol. Microbiol.* **14**, 115-121) were transformed into *S. aureus* OS2 (*spa:ermC*, r^-) (Schneewind, O., Model, P. and Fischetti, V. A. (1992) *Cell* **70**, 267-281) and have been described previously. Staphylococci were generally grown in tryptic soy broth or agar. All chemicals were purchased from Sigma unless indicated otherwise.

Characterization of Cell Wall Sorting Intermediates

S. aureus OS2 (pSeb-Spa490-524) was grown overnight in CDM (van de Rijn, I. and Kessler, R. E. (1980) *Infect. Immun.* **27**, 444-448) (Jeol BioSciences) supplemented with chloramphenicol (10 mg/ml), diluted 1:10 into minimal medium and grown with shaking at 37°C until A_{600} 0.6. Cells were labeled with 100 mCi of [^{35}S]-Promix (Amersham) for 1 minute. Labeling was quenched by the addition of an excess non-radioactive amino acid [50 ml chase (100 mg/ml casamino acids, 20 mg/ml methionine and cysteine)]. At timed intervals after the addition of the chase, 0, 1, 3, and 10 minutes, 250 ml aliquots were removed and protein was precipitated by the addition of 250 ml 10% TCA. The precipitate was sedimented by centrifugation 15,000 x *g* for 10 min, washed with 1 ml acetone and dried. Samples were suspended in 1 ml of 0.5 M Tris-HCl, pH 6.8 and staphylococcal peptidoglycan was digested by adding 50 ml lysostaphin (Schindler, C. A. and Schuhardt, V. T. (1964) *Proc. Natl. Acad. Sci. USA* **51**, 414-421) (100 mg, AMBI Pharmaceuticals) and incubating for 1 hour at 37°C. Proteins were again precipitated with TCA, washed with acetone and subjected to immunoprecipitation with a-Seb followed by SDS-PAGE and PhosphorImager analysis. To characterize the P1 and P2 precursors, 1 ml of culture was either incubated with 5 mM sodium azide for 5 min prior to labeling or 5 mM MTSET was added 15 seconds after the beginning of the pulse.

Antibiotic Inhibition of Cell Wall Sorting

Overnight cultures of *S. aureus* OS2 (pSeb-Spa490-524) grown in CDM were diluted into fresh minimal medium and incubated for until A_{600} 0.3. Cultures were then treated with either penicillin (10 mg/ml), vancomycin (10 mg/ml), moenomycin (10 mg/ml) or left untreated. A 0.5 ml culture sample was removed for pulse labeling with 100 mCi of [^{35}S]-Promix (Amersham) for 5 minutes. Labeling was quenched and proteins precipitated by the addition of 0.5 ml 10% TCA. The precipitate was collected by centrifugation, washed in acetone and dried under vacuum. The pellets were suspended in 1 ml 0.5 M Tris-HCl, pH 7.0, 50 ml lysostaphin (100 mg/ml, AMBI Pharmaceuticals) added and the staphylococcal cell wall digested

by incubating for 1 hour at 37°C. Proteins were precipitated with TCA, washed in acetone, dried and solubilized in 50 ml 0.5 M Tris-HCl, pH 7.5, 4% SDS and boiled for 10 min. Aliquots of solubilized surface protein were immunoprecipitated with a-Seb followed by SDS-PAGE and PhosphorImager analysis.

Peptidoglycan Synthesis Measurements

Staphylococci were grown in the presence or absence of antibiotics as described above. At 30 min intervals, 0.5 ml culture samples were withdrawn and labeled with either 50 mCi [^{3}H]lysine or 50 mCi [^{3}H]leucine for 20 min (Boothby, D., Daneo-Moore, L. and Shockman, G. D. (1971) *Anal. Biochem.* **44**, 645-653). All labeling was quenched by the addition of 0.5 ml 20% TCA. Samples were heated to 96°C for 30 min, cooled to room temperature and pipetted onto glass fiber filters. The filters were placed into a holder and washed under vacuum suction with 25 ml 75% ethanol and 2 ml 50 mM Tris-HCl, pH 7.8. After incubation in 5 ml pronase solution (50 mM Tris-HCl, pH 7.8, 1 mg/ml pronase) at 30°C for 30 min, filters were washed again with 4 ml of distilled water and 4 ml ethanol. The amount of radioactivity retained by the filter was determined by scintillation counting (Boothby, D., Daneo-Moore, L. and Shockman, G. D. (1971) *Anal. Biochem.* **44**, 645-653).

Chemical Inhibitors of the Sorting Reaction

S. aureus OS2 (pSeb-Spa490-524) was grown overnight in CDM supplemented with chloramphenicol (10 mg/ml), diluted 1:10 into minimal medium and grown with shaking at 37°C until A_{600} 0.6. Cells were labeled with 100 mCi of [^{35}S]-Promix (Amersham) for 5 minutes. Chemicals were added to a final concentration of 5 mM 15 seconds after the beginning of the pulse. All labeling was quenched by adding TCA to 10%. Precipitated cells and proteins were collected by centrifugation, washed in acetone and and the staphylococcal cell wall digested with lysostaphin as described above. The digests were again precipitated with TCA, immunoprecipitated with a-Seb followed by SDS-PAGE and PhosphorImager analysis.

Cell Wall Sorting in Staphylococcal Protoplasts

Overnight cultures of *S. aureus* OS2 (pSeb-Cws-BlaZ) or *S. aureus* OS2 (pSeb-Cws$_{DLPXTG}$-BlaZ) grown in CDM were diluted 1:10 into minimal medium and grown with shaking at 37°C until A_{600} 0.6. One ml of culture was pulse-labeled with 100 mCi of [^{35}S]-Promix (Amersham) for 2 minutes and labeling was quenched by the addition of 50 ml chase solution. Culture aliquots (0.5 ml) were removed for TCA precipitation either during the pulse or 20 min after the addition of chase. Another culture aliquot was first converted to protoplasts and then subjected to labeling. The cells were sedimented by centrifugation at 15,000 x*g* for 5 min and suspended in 1 ml 50 mM Tris-HCl, 0.4 M sucrose, 10 mM $MgCl_2$, pH 7.5. The cell wall was digested with lysostaphin (100 mg) for 30 min at 37°C. The protoplasts were labeled with 100 mCi of [^{35}S]-Promix (Amersham) for 2 minutes and labeling quenched by the addition of 50 ml chase solution. For sedimentation analysis, pulse-labeled staphylococci were centrifuged at 15,000 x*g* for 10 min to separate soluble surface protein from those that were bound to protoplasts. All samples were precipitated with TCA, washed in acetone and suspended in 50 ml 4% SDS, 0.5 M Tris-HCl pH 7.5 with boiling for 10 min. Aliquots of solubilized surface protein precursor and anchored products were immunoprecipitated with a-Seb and a-BlaZ, subjected to SDS-PAGE and PhosphorImager analysis.

EXAMPLE 3

Purification and Characterization of Sortase-Transpeptidase

To examine whether staphylococcal sortase captures surface proteins after their cleavage at the LPXTG motif as acyl-enzyme intermediates, the proposed acyl-enzyme intermediates between surface protein and sortase were treated by hydroxylaminolysis (P. Lawrence and J. L. Strominger, *J. Biol. Chem.* **245**, 3653 (1970); J. W. Kozarich, N. Tokuzo, E. Willoughby, J. L. Strominger, *J. Biol. Chem.* **252**, 7525 (1977)). In this model, the sulfhydryl of sortase may function as a nucleophile at the peptide bond between threonine and glycine, thereby forming a thioester with the carboxyl of threonine and releasing the amino of glycine (Figure 8A). Lipmann first used hydroxylamine to demonstrate the existence of acyl-enzyme intermediates as this strong nucleophile attacks thioester to form hydroxamate with carboxyl, thereby regenerating enzyme sulfhydryl (F. Lipmann and L. C. Tuttle, *J. Biol. Chem.* **161**, 415 (1945)).

Hydroxylaminolysis of Surface Proteins

Hydroxylaminolysis of surface proteins was examined by pulse-labeling staphylococci with [^{35}S]methionine in either the presence or absence of 0.2 M NH_2OH. Cultures were labeled with [^{35}S]methionine and divided into two aliquots, each of which was precipitated with 5% TCA. One sample was boiled in hot SDS, whereas the other was first treated with lysostaphin to release all anchored surface protein, and then boiled in hot SDS. Surface protein (SEB-SPA490-524) of mock treated staphylococci was insoluble in hot SDS (3.8%) unless the peptidoglycan had been digested with lysostaphin prior to boiling in SDS (100%)(Figure 12A). Addition of 0.2 M NH_2OH caused 25.3% of all labeled SEB-SPA490-524 to be released into the extra-cellular medium and to be soluble in hot SDS. This phenomenon was not strain specific as *S. aureus* OS2 and *S. aureus* BB270 displayed similar amounts of surface protein hydroxylaminolysis.

If the solubility of surface proteins in hot SDS is caused by hydroxylaminolysis of acyl-enzyme intermediates, addition of NH_2OH after the pulse labeling of staphylococci should not release SEB-SPA490-524 as this polypeptide is rapidly anchored to the cell wall. Addition of NH_2OH either before or during the pulse with [^{35}S]methionine released surface proteins into the extra-cellular medium (16.9% and 12.7%, respectively) (Figure 12B). Very little SDS-soluble SEB-SPA490-524 was detected when NH_2OH was added after the pulse (4%). Increasing the amount of NH_2OH prior to pulse-labeling resulted in increased amounts of released surface proteins (Figure 12C).

Characterization of NH_2OH-released Surface Proteins

Hydroxylaminolysis of sortase acyl-intermediates should result in the formation of surface protein hydroxamate at the threonine of the LPXTG motif. To characterize NH_2OH-released surface protein, staphylococci (10^{13} cfu) expressing the surface protein SEB-MH_6-CWS (H. Ton-That, K. F. Faull, O. Schneewind, *J. Biol. Chem.* **272**, 22285 (1997)) were incubated in the presence or absence of 0.1 M NH_2OH. Samples were centrifuged to sediment bacteria and SEB-MH_6-CWS was purified from the supernatant by affinity chromatography and analyzed on Coomassie-stained SDS-PAGE. Treatment with 0.1 M NH_2OH caused the release of SEB-MH_6-CWS by *S. aureus* strains OS2 and BB270 (Figure 13A). SEB-MH_6-CWS purified from strain BB270 was cleaved at methionine with cyanogen bromide. COOH-terminal peptides bearing anchor structures were purified by affinity chromatography and analyzed by rpHPLC (H. Ton-That, K. F. Faull, O. Schneewind, *J. Biol. Chem.* **272**, 22285 (1997)). The chromatogram of anchor peptides released from mock treated bacteria revealed a major absorbance peak at 29% CH_3CN (Figure 13B). The sample was subjected to electrospray-ionization mass spectrometry (ESI-MS) and a compound with an average mass of 2236 Da was detected. This measurement is consistent with the structure of anchor peptide linked to a branched cell wall tetrapeptide [L-Ala-D-iGln-L-Lys(NH_2-H_6AQALPET-Gly5)-D-Ala,

predicted mass 2235]. This surface protein species is not linked to the glycan strands of the staphylococcal cell wall and is therefore released into the culture medium. The chromatogram of anchor peptides released by treatment with 0.1 M NH_2OH revealed a major absorbance peak at 32% CH_3CN (Figure 13C). ESI-MS identified a compound with the average mass of 1548 Da. When subjected to Edman degradation, the peptide sequence NH_2-H_6AQALPET* was obtained, in which the thirteenth cleavage cycle released a phenylthiohydantoin moiety of unknown structure. The predicted mass of NH_2-H_6AQALPET> (T> indicates threonine hydroxamate) is 1565 Da, 17 Da more than the observed mass of 1548 Da. Fractions of both chromatograms were scanned by rpHPLC for the presence of ion signals with an average mass of 1548, 1565 or 2236. rpHPLC fractions of anchor peptides from mock-treated cultures contained the compound with mass 2236, however no ions of the predicted mass 1548 or 1565 were detected. In contrast, rpHPLC fractions collected from anchor peptides of NH_2OH-treated staphylococci harbored compounds with an average mass of 1548 Da (NH_2-H_6AQALPET*, 32% CH_3CN) and 1565 Da (NH_2-H_6AQALPET>, 31% CH_3CN), but not the anchor peptide of 2235 Da. Thus, treatment with 0.1 M NH_2OH released surface protein from staphylococci as a hydroxamate of the threonine within the LPXTG motif, suggesting that sortase forms an acyl-enzyme intermediate with cleaved surface protein. The peptide NH_2-H_6AQALPET> appears to be unstable during our purification, thereby generating NH_2-H_6AQALPET* with a loss of 17 Da at the threonine hydroxmate.

Analysis of Sortase Hydroxylaminolysis Activity *In Vitro* in the Presence of NH_2OH

If NH_2OH can release surface protein from staphylococci *in vivo*, sortase may catalyze the cleavage of LPXTG motif bearing peptides in the presence of NH_2OH *in vitro*. Fluoresence of the EDANS fluorophore within the peptide DABCYL-QALPETGEE-EDANS is quenched by the close proximity of DABCYL (G. T. Wang, E. Matayoshi, H. J. Huffaker, G. A. Krafft, *Tetrahedon Lett.* **31**, 6493 (1990)). When the peptide is cleaved and the fluorophore separated from DABCYL, an increase in fluorescence is observed (E. D. Matayoshi, G. T. Wang,

G. A. Krafft, J. Erickson, *Science* **247**, 954 (1989)). Incubation of the LPXTG peptide with crude staphylococcal extracts caused only a small increase in fluorescence. However, the addition of 0.1 M NH_2OH to staphylococcal extracts resulted in a forty fold increase in fluorescence intensity (Figure 14). This activity appears to be specific for sortase as it can be inhibited by pre-incubation of staphylococcal extracts with methanethiosulfonate (MTSET) (D. J. Smith, E. T. Maggio, G. L. Kenyon, *Biochemistry* **14**, 764 (1975), a known inhibitor of the sorting reaction. These results suggest that sortase catalyzes the hydroxylaminolysis of LPXTG peptide *in vitro*. Thus, surface protein is cleaved between the threonine and the glycine of the LPXTG motif, resulting in the formation of a NH_2OH-sensitive thioester linkage between the carboxyl of threonine and the active site sulfhydryl of sortase. *In vivo*, the acyl-enzyme intermediate is resolved by a nucleophilic attack of the amino within the pentaglycine crossbridge. Recent observations suggest that the pentaglycine crossbridge of the lipid II precursor functions as a nucleophile for the sorting reaction. We show here that hydroxylamine can subsitute for pentaglycine both *in vivo* and *in vitro*.

Purification and Characterization of Sortase

When expressed in *E. coli* and analyzed by centrifugation of crude lysates, the staphylococcal SrtA protein sedimented with membranes. To obtain a soluble enzyme and to examine its properties, the NH_2-terminal membrane anchor segment of SrtA was replaced with a six histidine tag ($SrtA_{DN}$). $SrtA_{DN}$ was expressed in *E. coli* XL-1Blue and purified by affinity chromatography from cleared lysates. When incubated with the LPXTG peptide and measured as an increase in fluorescence, $SrtA_{DN}$ catalyzed cleavage of the substrate. Addition of 0.2 M NH_2OH to this reaction resulted in an increase in fluorescence, indicating that cleavage of the LPXTG peptide occurred more efficiently. Hydroxylaminolysis of LPXTG peptide was dependent on the sulfhydryl of $SrtA_{DN}$ as pre-incubation with MTSET abolished all enzymatic activity. Methanethiosulfonate forms disulfide with sulfhydryl (D. J. Smith, E. T. Maggio, G. L. Kenyon, *Biochemistry* **14**, 764 (1975); M. H. Akabas and A. Karlin, *Biochemistry* **34**, 12496

(1995)) which can be reversed by reducing reagents such as dithiothreitol (DTT) (R. Pathak, T. L. Hendrickson, B. Imperiali, *Biochemistry* **34**, 4179 (1995)). MTSET-inactivated $SrtA_{DN}$ was incubated in the presence of 10 mM DTT, which restored 80% of LPXTG peptide cleavage activity. The availability of purified, soluble sortase ($SrtA_{DN}$)and an *in vitro* assay for the hydroxylaminolysis of LPXTG peptide should allow the screening for compounds that interfere with the anchoring of surface protein in Gram-positive bacteria. Such compounds may be useful for the therapy of human infections with Gram-positive bacteria that have gained resistance to all known antibiotics.

Materials and Methods

Pulse-Chase Screen of Hydroxylaminolysis of surface proteins

Staphylococci were grown in minimal medium unil OD_{600} 0.6 and pulse-labeled with 100 µCi Pro-Mix ([^{35}S] methionine and cysteine) for 1 min. Incorporation of radio-label into polypeptides was quenched by the addition of 50 µl chase solution (100 mg/ml casamino acids, 20 mg/ml methionine and cysteine) and incubation was continued at 37°C for 5 min. Two 0.5 ml aliquots of labeled culture were each precipitated with 0.5 ml 10% TCA, washed in acetone and dried under vacuum. One sample was suspended in 50 µl 0.5 M tris, 4% SDS and boiled. The other sample was first suspended in 1 ml 0.5 M Tris pH 7.0 and the cell wall digested for 1 hour at 37°C by adding 50 µl 2 mg/ml lysostaphin. The sample was precipitated with 75 µl 100% TCA, washed in acetone, dried and then boiled in SDS. Aliquots were subjected to immunoprecipitation with a-SEB and analyzed after SDS-PAGE on PhosphorImager.

Purification of NH_2OH Surface Proteins

Staphylococci (10^{13} cells) were incubated in 200 ml 50 mM Tris-HCl, pH 7.0 with or without 0.1 M NH_2OH for 60 min. Sampes were centrifuged at 10,000 x*g* for 15 min and the supernatants applied to 1 ml Ni-NTA column, pre-equilibrated with column buffer (CB, 50 mM Tris-HCl, 150 mM NaCl, pH 7.5). The column was washed first with 20 ml CB and 20

ml CB containing 10% glycerol and eluted with 4 ml of column buffer and 0.5 imidazol. Aliquots were mixed with sample buffer and separated on SDS-PAGE. The eluate was precipitated with TFA (10%), washed in acetone and dried under vacuum. The sample was suspended in 600 µl 70% formic acid and, after addition of a crystal of cyanogen bromide, incubated overnight. Cleaved peptides were repeatedly dried and suspended in water to evaporate cyanogen bromde, solubilized in 1 ml buffer A and subjected to affinity chromatography as previusly described. Peptides were eluted in 4 ml of 6 M guanidine-hydrochloride, 0.2 M acetic acid, desalted over C18 cartridge and dried. Pellets were solubilized in 50 µl buffer B (8 M urea, 50 mM phosphate, 10 mM Tris-HCl, pH 7.3) and subjected to rpHPLC on C18 column (Hypersil, Keystone Scientific) with a linear gradient from 1%-99% CH_3CN in 0.1% TFA in 90 minutes. MALDI-MS and ESI-MS was performed as described (H. Ton-That, K.F. Faull, O. Schneewind (1997) *J. Biol. Chem.* 272:22285-22292).

Identification of peptide structure by Mass Spectrometry

The structure of the peptides with mass 1548 and 1565 was determined by tandem mass spectrometry, MS/MS using the parent ions. Collisionally induced dissociation of the parent ions produced daughter ion spectra consistent with compound structures NH_2-H_6AQALPET> (T> is threonine hydroxamate, predicted compound mass 1565) and NH_2-H_6AQALPET* (T* represents a loss of 17 Da of threonine hydroxamate; the structure of this residue is unknown).

Assay of Sortase activity by Fluorescent Assay

Reactions were assembled in a volume of 120 µl containing 50 mM Tris-HCl, 150 mM NaCl, pH 7.5. The concentration of LPXTG peptide substrate (DABCYL-QALPETGEE-EDANS) was 10 µM, of MTSET 5 mM, of NH_2OH 0.2 M. Staphylococcal cell extracts were obtained by subjecting 1013 cells to disrubtion in a bead beater instrument. The crude extract was subjected to slow speed centrifugation at 3,000 x*g* for 15 min to remove beads and intact cells. A 10 µl aliquot of the supernatant, containing approximately 50 mg/ml protein, was used

as enzyme preparation. Incubations were carried out for 1 hour at 37°C, followed by centrifugation of the sample at 15,000 x*g* for 5 min. The supernatant was subjected to analysis in a fluorimeter using 395 nm for excitation and 495 nm for recordings.

Purification of Sortase by Addition of Histidine Tag

The primers orf6N-ds-B (5'-AAAGGATCCAAACCACATATCGATAATTATC-3') and orf6C-dT-B (5'-AAAGGATCCTTTGACTTCTGTAGCTACAAAG-3') were used to PCR amplify the *srtA* sequence from the chromosome of *S. aureus* OS2. The DNA fragment was cut with *Bam*HI, inserted into pQE16 (Qiagen) cut *Bam*HI to generate pHTT5, transformed into *E. coli* XL-1 Blue and selected on Luria broth with ampicillin (100 µg/ml). *E. coli* XL-1 Blue (pHTT5) (10^{12} cells) were suspended in 30 ml C buffer (50 mM Bis-Tris-HCl, 150 mM NaCl, 10% glycerol, pH 7.2) and lysed by one passage through a French pressure cell at 14,000 psi. The extract was centrifuged at 29,000 x*g* for 30 min and the supernatant applied to 1 ml Ni-NTA resin, pre-equilibrated with C buffer. The column was washed with 40 ml C buffer and $SrtA_{DN}$ protein was eluted in 4 ml C buffer with 0.5 M imidazol at a concentration of 30 µg/µl.

Reactions were assembled in a volume of 260 µl containing 50 mM Hepes buffer, 150 mM NaCl, pH 7.5 and as indicated 5 µM $SrtA_{DN}$ in 50 mM BisTris, pH 7.5, 10 µM LPXTG peptide (DABCYL-QALPETGEE-EDANS), 10 µM TGXLP peptide (DABCYL-QATGELPEE-EDANS), 5 mM MTSET, 0.2 M NH_2OH, 5 mM pHMB or 10 mM DTT. Incubations were carried out for 1 hour at 37°C. Samples were analyzed in a fluorimeter using 395 nm for excitation and 495 nm for recordings.

ADVANTAGES OF THE PRESENT INVENTION

In isolating and characterizing the gene for the *S. aureus* sortase-transamidase enzyme, we have determined the existence of a new site for antibiotic action that can be used to screen new antibiotics active against Gram-positive pathogens, such as *Staphylococcus*, *Actinomyces*, *Mycobacterium*, *Streptococcus*, *Bacillus*, and other medically important Gram-positive pathogens increasingly resistant to conventional antibiotics. The availability of substantially purified *S. aureus* sortase-transamidase enzyme provides a method of screening compounds for inhibition of the enzyme.

The purified sortase-transamidase enzyme of the present invention also yields a method of surface display of peptides and proteins that has advantages over phage display, as well as providing methods for producing vaccines against a large variety of antigens that can be covalently bound to the surfaces of Gram-positive bacteria.

Although the present invention has been described with considerable detail, with reference to certain preferred versions thereof, other versions and embodiments are possible. Therefore, the scope of the invention is determined by the following claims.

TABLE I

Inhibition of the sorting reaction by methanethiosulfonates and organic mercurial

The sorting reaction was measured as the ratio between the amount of pulse-labeled Seb-Spa490-524 P2 precursor [P2] and the mature, anchored species processed at the LPXTG motif [M].

Compound (5 mM)	[P2]/[M]
[2-(trimethylammonium)ethyl]methanethiosulfonate (MTSET)	23.14 ± 0.06[a]
(2-sulfonatoethyl)methanethiosulfonate (MTSES)	1.61± 0.03
p-hydroxymercuribenzoic acid (pHMB)	1.51 ± 0.04
phenylmethylsulfonylfluoride (PMSF)	0.16 ± 0.05
N-ethylmaleimide	0.16 ± 0.05
iodoacetamide	0.12 ± 0.01
iodoacetic acid	0.13 ± 0.02
2-mercaptoethanol	0.15 ± 0.04
dithiothreitol (DTT)	0.13 ± 0.03
zinc chloride ($ZnCl_2$)	0.32 ± 0.02
calcium chloride ($CaCl_2$)	0.06 ± 0.05
magnesium chloride ($MgCl_2$)	0.13 ± 0.01
ethylenediaminetetraacetic acid (EDTA)	0.31 ± 0.04
mock treated	0.15 ± 0.02

[a]Data represent an average of three measurements. The standard deviation is indicated as ±.

TABLE II

Antibiotic inhibition of cell wall synthesis and the effect on cell wall sorting

The cell wall sorting reaction was measured as the ratio between the amount of pulse-labeled Seb-Cws-BlaZ precursor [P] and the mature, anchored species processed at the LPXTG motif [C]. Cell wall synthesis was measured as the ratio between the amount of [^{3}H]lysine and that of [^{3}H]leucine incorporated into the acid precipitable, pronase resitant peptidoglycan. The data are presented as percent inhibition.

Compound	[P2]/[M][a]	fold inhibition of cell wall synthesis[a]
vancomycin (10 μg/ml)	0.47± 0.04	9.5
moenomycin (10 μg/ml)	0.24± 0.04	1.6
penicillin (10 μg/ml)	0.10± 0.01	3.3
untreated	0.15± 0.02	-

[a]Data were collected from cultures that were grown for 60 min in the presence of antibiotics.